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(54) Rinse-off water-in-oil-in-water compositions

(57) Rinse-off, water-in-oil-in-water multiple emulsion compositions are disclosed. The multiple emulsion compositions comprise an external aqueous phase optionally incorporating an emulsifier and/or a second topically-active compound. The internal phase comprises a primary water-in-oil emulsion, wherein the primary emulsion comprises a first topically-active compound, a surfactant phase, an oil phase, and water.

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Description**FIELD OF THE INVENTION**

5 The present invention is directed to rinse-off water-in-oil-in-water (W_1-O-W_2) multiple emulsion compositions comprising: (1) a primary water-in-oil (W_1/O) emulsion as the internal phase and (2) an external aqueous phase. The primary W_1/O emulsion comprises (a) a first topically-active compound; (b) a surfactant or blend of surfactants; (c) an oil phase comprising a silicone compound, a hydrocarbon compound or a mixture thereof; and (d) water. The external aqueous phase comprises water, and preferably further comprises (a) an emulsifier, such as a nonionic surfactant or a surfactant system capable of forming stabilizing liquid crystals and/or (b) a second topically-active compound. In use, a W_1-O-W_2 multiple emulsion composition is applied to the skin or hair to deliver the second topically-active compound. The external aqueous phase of the composition then is rinsed from the skin or hair. The primary (W_1/O) emulsion deposits on and remains on the hair or skin during rinsing. After evaporation of the oil phase or after rupture of the W_1/O emulsion by friction (e.g., rubbing), the first topically-effective compound remains on the skin or hair to perform its intended function.

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BACKGROUND OF THE INVENTION

An ideal composition for delivery of a topically-active compound to skin or hair delivers the topically-active compound such that it adheres to the skin or hair, while topically-inactive ingredients evaporate or are rinsed from the application site. Delivery of a water-insoluble topically-active compound is facilitated because the topically-active compound is not removed from the application site during the rinsing step. The difficulty with this type of composition resides in incorporating the water-insoluble topically-active compound into an aqueous composition. This difficulty is overcome by forming emulsions. Therefore, topically-active compounds, such as a hair conditioner, a hair dye, a skin care compound or a topical medicament, conventionally have been delivered from either oil-in-water emulsions or water-in-oil emulsions.

20 With respect to water-soluble and water-dispersible topically-active compounds (hereinafter collectively termed "water-soluble topically-active compounds"), it is easy to incorporate the topically-active compound into an aqueous rinse-off composition. However, a substantial amount, or all, of the water-soluble compound is rinsed from the treated skin or hair. The water-soluble compound therefore cannot perform its intended function and is wasted. Accordingly, rinse-off compositions often incorporate an excess of the water-soluble topically-active compound, such that a sufficient amount of the compound remains on the skin or hair after rinsing to perform its intended function. The need to overformulate is economically disadvantageous.

25 For example, water-soluble dyes are used in hair products because these dyes are semi-permanent, i.e., temporary dyes that do not dye the hair for extended time periods or through numerous shampooings. However, water-soluble hair dyes are difficult to apply because a substantial amount of applied dye is removed from the hair during the rinsing operation. Compositions incorporating water-soluble dyes also stain the hands of the applicator and various articles in the work area, such as sinks and countertops, both during product application and rinsing of the product from the hair.

30 Similarly, two-in-one products, like shampoo-conditioner compositions, have been difficult to formulate not only because of the water solubility of many quaternary ammonium compounds, but also because of the well-known incompatibility between an anionic surfactant and a quaternary ammonium compound. Formulators either have avoided this problem by sequentially contacting the hair first with an anionic surfactant, then a quaternary ammonium compound from different compositions, or have overcome this problem by using silicone conditioners in shampoo-conditioners. However, silicones and similar conditioners are water insoluble and an esthetically-unpleasant residue builds up on the hair after repeated applications. In addition, quaternary ammonium compounds impart hair conditioning properties different from a silicone, and therefore it is desirable to condition the hair with quaternary ammonium compounds in place of, or in conjunction with, a silicone.

35 Investigators therefore have searched for rinse-off, topically-effective compositions that overcome the above-described disadvantages, and that have consumer-acceptable stability and esthetics. A rinse-off, topically-effective composition however often is difficult to formulate because of the water solubility of the desired topically-active compound.

40 Many aqueous compositions are designed to be rinsed from the skin or hair after application. These include hair conditioners, hair shampoos, skin cleaners, facial scrubs, permanent wave lotions and temporary hair dyes. For many rinse-off products, it is desirable to leave one or more topically-active compounds on the skin or hair to perform an intended function. When the topically-active compound is water soluble, it is difficult, and often impossible, for a sufficient amount of the compound to remain on the skin or hair to be efficacious. Investigators therefore have directed their attention to compositions designed to be rinsed from the skin or hair after application and that are capable of depositing efficacious amounts of water-soluble, topically-active compounds to hair or skin.

45 One type of composition that has been investigated are multiple emulsions. Multiple emulsions are complex emulsions of emulsions formed by dispersing droplets in a continuous phase, wherein the droplets themselves contain smaller droplets of a liquid similar to the external continuous phase. One type of multiple emulsion is a water-in-oil-in-water emulsion (W_1-O-W_2), wherein a primary water-in-oil (W_1/O) emulsion is dispersed in an external aqueous phase (W_2).

Multiple emulsions also are known as liquid membrane systems because, in the case of a W₁-O-W₂ emulsion, an organic membrane, film or layer separates the inner water droplets from the external aqueous phase.

W₁-O-W₂ multiple emulsion compositions have been used in the pharmaceutical industry as vaccine adjuvants and as sustained release and parental drug delivery systems. However, the low stability of W₁-O-W₂ multiple emulsion compositions has limited their widespread use.

Multiple emulsion compositions also have been used in cosmetics and skin care products. W₁-O-W₂ multiple emulsions however were difficult to prepare and were unstable at elevated temperatures. In addition, various active ingredients had a tendency to further destabilize the multiple emulsions, for example causing leakage between the external aqueous phase and internal aqueous phase.

W₁-O-W₂ multiple emulsion compositions have been disclosed in numerous patents and publications. For example, M.C. Taelman et al., "Multiple Emulsions in Cosmetics", Publication RP112/94E, March, 1994, ICI Europe Limited, Everberg, Belgium, discloses multiple emulsions in general, including methods of manufacturing multiple emulsions. S. Matsumoto et al., *J. Dispersion Science and Technology*, 10 (1989), pp. 455-482, discloses the use of multiple emulsions in the food, drug and cosmetic industries.

Other publications which disclose multiple W₁-O-W₂ emulsions are:

S.S. Davis et al., *Encyclopedia of Emulsion Technology*, Vol. 2; P. Becher, Ed.; Marcel Dekker, Inc., New York, NY (1985), pp. 159-238;

Y. Sela et al., *Colloids and Surfaces A: Physicochemical and Engineering Aspects*, 83 (1994), pp. 143-150;

K. Oza et al., *J. Dispersion Science and Technology*, 10(2) (1985), pp. 163-185;

J. Zatz et al., *J. Soc. Cosmet. Chem.*, 39 (May/June 1988), pp. 211-222;

C. Fox, *Cosmetics and Toiletries*, 101 (Nov. 1986), pp. 101-106 and 109-112;

M. deLuca et al., *Cosmetics and Toiletries*, 105 (Nov. 1990), pp. 65-66 and 69;

T. Tadros, "Application of Multiple Emulsions in Cosmetics", ICI Surfactant publication RP56/92E (1992); and

C. Prybilsky et al., *Cosmetics and Toiletries*, 106 (Nov. 1991), pp. 97-100.

Various patents disclose multiple emulsion compositions used in cosmetic compositions. For example, WO 94/01073 discloses gelled multiple emulsion compositions for cosmetic use. Other patents include WO 94/02120; WO 93/00160; WO 92/18227; JP 58 183 611; U.S. Patent No. 5,306,498; and U.S. Patent No. 4,988,456.

U.S. Patent Nos. 5,277,899 and 4,963,348, and WO 93/07848, disclose conditioners that have set retention properties. The disclosed compositions are oil-in-water emulsions (O/W), as opposed to the present W₁/O primary emulsions

and the W₁-O-W₂ multiple emulsion compositions.

Although various patents and publications disclose W₁-O-W₂ multiple emulsion compositions for cosmetic use, those compositions are designed as leave-on products, i.e., products that are applied to, and not subsequently rinsed from, the skin or hair, and that are allowed to remain in contact with the skin or hair. The present invention therefore is directed to W₁-O-W₂ multiple emulsion compositions which are capable of depositing water-soluble, topically-active compounds onto hair or skin from rinse-off compositions, such as hair conditioners, hair shampoos, or skin cleaners.

SUMMARY OF THE INVENTION

The present invention is directed to rinse-off, topically-effective W₁-O-W₂ multiple emulsion compositions having improved efficacy, and to methods of using the W₁-O-W₂ multiple emulsion compositions. More particularly, the present invention is directed to a W₁-O-W₂ multiple emulsion composition comprising a continuous external aqueous phase (W₂) and a primary water-in-oil (W₁/O) emulsion as the dispersed internal phase.

The primary W₁/O emulsion comprises a first topically-active compound, like a hair conditioner, a hair dye, a hair fixative, a skin care compound or a topical medicament; a surfactant phase comprising silicon-free surfactant or surfactant blend having an HLB value of about 10 or less, an oil-soluble silicon-based surfactant, an oil-soluble polymeric surfactant or a mixture thereof; an oil phase comprising a silicone compound or a hydrocarbon compound; and water. The external phase comprises water, and preferably a second topically-active compound and/or an emulsifier, such as a nonionic surfactant having an HLB value of greater than about 10 or a surfactant system capable of forming stabilizing liquid crystals.

The "HLB value", or hydrophobic-lipophilic balance value, of a surfactant is a term well-known to those skilled in the art. The HLB value is related to the solubility of the surfactant, wherein a surfactant with a low HLB value, e.g., about 10 or less, tends to be oil soluble and a surfactant with a high HLB value, e.g., greater than about 10, tends to be water soluble.

In particular, the W₁-O-W₂ multiple emulsion compositions comprise:

- (a) about 40% to about 99% by weight of the composition of an external aqueous phase (W₂); and
- (b) about 1% to about 60% by weight of the composition of a primary W₁/O emulsion.

The primary W₁/O emulsion comprises:

- (a) about 1% to about 95% by weight of the primary emulsion of an aqueous phase comprising (i) a first topically-active compound and (ii) water;
- (b) about 0.5% to about 95% by weight of the primary emulsion of an oil phase comprising a volatile silicone compound, a nonvolatile silicone compound, a volatile hydrocarbon compound, a nonvolatile hydrocarbon compound, or a mixture thereof; and
- (c) about 0.1% to about 20% by weight of the primary emulsion of a surfactant phase comprising a silicon-free surfactant or surfactant blend having an HLB value of about 10 or less, an oil-soluble silicon-based surfactant, an oil-soluble polymeric surfactant, or a mixture thereof.

10 The external aqueous phase (W_2) optionally includes (a) an emulsifier, such as a nonionic surfactant having an HLB value of greater than about 10 or a surfactant system capable of forming stabilizing liquid crystals and/or (b) a second topically-active compound which can perform the same or a different function from the first topically-active compound. The external aqueous phase is free of a polymeric gelling agent, such as a polyacrylic acid.

15 A W_1 -O- W_2 multiple emulsion composition of the present invention can be a liquid composition having a viscosity of about 1 cps to about 15,000 cps (centipoise), and preferably about 100 to about 10,000 cps. To achieve the full advantage of the present invention, a liquid topically-effective composition has a viscosity of about 1,000 to about 9,000 cps.

20 The W_1 -O- W_2 multiple emulsion also can be cream-like and have a viscosity of about 50,000 to about 1,200,000 cps, and preferably about 100,000 to about 1,000,000 cps. A W_1 -O- W_2 composition having a cream-like consistency is free of a polymeric gelling agent in the external phase.

25 The W_1 -O- W_2 multiple emulsion compositions of the present invention are rinse-off compositions, which are designed for application to the skin or hair, followed by rinsing from the skin or hair. The second topically-active compound, if present in the external aqueous phase, performs its intended function during and after topical application of the composition to the skin or hair, then is rinsed from the skin or hair. The droplets of the primary W_1 /O emulsion are of sufficient size, i.e., a range of about 5 to about 1000 μ (microns), and preferably about 10 to about 500 μ , in diameter, to resist removal from the skin or hair during the rinsing step. The droplets of the internal W_1 phase of the W_1 /O primary emulsion range in size from about 1 to about 75 μ .

30 The external oil phase of the primary W_1 /O emulsion then evaporates during drying of the skin or hair to release the first topically-active compound present in the internal aqueous phase of the primary emulsion. Alternatively, the internal aqueous phase is released by rubbing the skin or hair to break the W_1 /O primary emulsion. The first topically-active compound then can contact the skin or hair to perform its intended function. In another embodiment, the primary W_1 /O emulsion is not broken, but is allowed to remain on the skin or hair as is. Typically, the first topically-active compound is not rinsed from the skin or hair.

35 In accordance with an important aspect of the present invention, a water-soluble (or water-dispersible), topically-active compound therefore is included in a rinse-off composition without wasting a substantial amount of the topically-active compound during the rinsing step.

40 Another important aspect of the present invention is the ability to treat the skin or hair with incompatible water-soluble, topically-active compounds using a single W_1 -O- W_2 multiple emulsion composition. By incorporating the first topically-active compound in the internal aqueous phase of the primary emulsion (W_1) and the second topically-active compound in the external aqueous phase of the composition (W_2), the oil phase (O) provides a barrier that separates the two incompatible topically-active compounds and precludes contact between the compounds. Incompatible topically-active compounds therefore can be applied to the hair or skin from a single composition, as opposed to applying the topically-active compounds individually from two different compositions.

45 For example, it is well known that an anionic compound, like an anionic surfactant, and a cationic compound, like a quaternary ammonium compound, are incompatible. However, these incompatible compounds can be included in the present W_1 -O- W_2 multiple emulsion compositions.

50 The anionic surfactant is present as the second topically-active compound in the external aqueous phase (W_2). Upon application to the hair, the anionic surfactant cleanses the hair, then is rinsed from the hair. The primary W_1 /O emulsion, which incorporates the quaternary ammonium compound (i.e., the first topically-active compound) in the aqueous phase, is not rinsed from, but remains on, the hair. An oil phase provides conditioning benefits to the wet hair, and evaporates during hair drying to release the quaternary ammonium compound, which is substantive to, and further conditions, the hair. Similarly, a cationic compound can be the second topically-active compound to condition the hair and an anionic hair dye can be the first topically-active compound to dye the hair.

55 In accordance with another important aspect of the present invention, the first topically-active compound is water soluble or water dispersible. The first topically-active compound therefore can be rinsed from the skin or the hair during a subsequent shampooing or rinsing to preclude an undesirable residue of first topically-active compound building up on the hair.

In another embodiment of the present invention, the first and second topically-active compounds are compatible, yet perform different functions. For example, the second topically-active compound can be a cleansing compound for

the skin or hair, and the first topically-active compound can be a topical medicament, a hair fixative, or a hair dye, for example, designed to remain on the skin or hair.

In another important embodiment, the external aqueous phase does not include a second topically-active compound. In this embodiment, the first topically-active compound in the primary W₁/O emulsion is precluded from contacting the skin or hair until desired. For example, present day water-soluble hair dye compositions stain the hands of the applicator, sinks and countertops. The present W₁-O-W₂ multiple emulsion compositions allow application of a water-soluble dye to hair from a rinse-off product, wherein the dye is released only after the composition is applied to the hair, and the hair is rinsed and dried. Staining of hands, sinks and countertops therefore is avoided. Composition economics also is enhanced because the water-soluble hair dye, a relatively expensive ingredient, is not rinsed from the hair and wasted.

In yet another important embodiment, a W₁-O-W₂ multiple emulsion composition of the present invention incorporates a cleansing agent as the second topically-active compound and an astringent salt as the first topically-active compound to form a stable and efficacious deodorant-cleanser composition. In other embodiments of the present invention, the first topically-active compound is a topically-effective drug or medicament; a topical anesthetic; a sunscreen agent; a skin care agent; a skin-soothing emollient or other topical cosmetic compound; a topical anti-inflammatory; and the like.

The first topically-active compound incorporated into the multiple emulsion compositions of the present invention preferably is water soluble and is incorporated into the aqueous phase of the primary W₁/O emulsion. As used herein, the term "water soluble" means "water soluble or water dispersible". However, water-insoluble topically-active compounds also can be incorporated into the oil phase of the primary W₁/O emulsion.

In a preferred embodiment, the W₁-O-W₂ multiple emulsion compositions comprise:

(A) about 50% to about 95% by weight of the composition of an external aqueous phase, said external aqueous phase comprising:

- (i) about 0.1% to about 30% by weight of the external aqueous phase of a second topically-active compound;
- (ii) about 0.1% to about 10% by weight of the external aqueous phase of an emulsifier; and
- (iii) water;

(B) about 5% to about 50% by weight of the composition of a primary water-in-oil emulsion, said primary emulsion comprising:

- (i) about 10% to about 95% by weight of the primary emulsion of an aqueous phase, said aqueous phase comprising (a) water and (b) a topically-effective amount of a first topically-active compound, typically about 0.1% to about 30% by weight based on the total weight of the primary emulsion;

- (ii) about 0.5% to about 80% by weight of the primary emulsion of an oil phase comprising a volatile silicone compound, a volatile hydrocarbon compound, or a mixture thereof; and

- (iii) about 0.1% to about 15% by weight of the primary emulsion of a surfactant phase comprising a silicon-free surfactant or surfactant blend having an HLB value of about 1 to about 7, an oil-soluble silicon-based surfactant, an oil-soluble polymeric surfactant or a mixture thereof.

Other aspects of the present invention include providing a phase stable, topically-effective composition, which is free of polymeric gelling agents in the external aqueous phase, for the administration of one or more water-soluble, topically-active compounds from a rinse-off product by providing a primary W₁/O emulsion having a droplet size of about 5 to about 1000 μ .

The present W₁-O-W₂ multiple emulsion compositions are stable over extended storage periods, can be essentially

nonstaining, effectively deliver one or more water-soluble, topically-active compounds to the skin and hair, and exhibit excellent esthetic and functional properties for consumer acceptance.

The above and other advantages and novel features of the present invention will become apparent from the following detailed description of the preferred embodiments.

50 DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

A topically-effective composition of the present invention is a stable, W₁-O-W₂ multiple emulsion comprising an external aqueous phase and an internal primary emulsion. The primary emulsion comprises a first water-soluble, topically-active compound, such as a hair conditioner, a hair fixative, a hair dye or a topical medicament; a surfactant phase

comprising a silicon-free surfactant or surfactant blend having an HLB value of about 10 or less, an oil-soluble silicon-based surfactant, an oil-soluble polymeric surfactant, or a mixture thereof; an oil phase comprising a volatile silicone compound, a nonvolatile silicone compound, a volatile hydrocarbon compound, a nonvolatile hydrocarbon compound, or a mixture thereof; and water.

The W₁-O-W₂ multiple emulsion composition is a rinse-off composition. The W₁-O-W₂ multiple emulsion composition can be a liquid having a viscosity in the range of about 1,000 to about 15,000 cps. The multiple emulsion composition also can be a cream having a viscosity of about 50,000 to about 1,200,000 cps. The external phase of the W₁-O-W₂ multiple emulsion composition is free of a polymeric gelling agent.

5 After application to the skin or hair, the external aqueous phase is rinsed from the skin or hair. The oil phase of the primary emulsion, if sufficiently volatile, evaporates during the skin or hair drying process to release the internal aqueous phase of the primary emulsion, thereby allowing the first topically-active compound to contact the skin or hair and perform its intended function. Alternatively, the internal aqueous phase is released after the rinsing step by rubbing the skin or hair to break the primary emulsion.

10 The water-in-oil-in-water multiple emulsion compositions of the present invention are designated herein as W₁-O-W₂ multiple emulsions. This designation indicates that an oil phase (O) separates an encapsulated, discontinuous aqueous phase (W₁) from a continuous aqueous phase (W₂). Conventionally, the W₁ and W₂ phases differ in compositional make-up, but the W₁ and W₂ phases can be identical.

15 In accordance with an important feature of the present invention, the liquid W₁-O-W₂ multiple emulsion compositions are stable compositions that effectively resist droplet coalescence over long storage periods. The W₁-O-W₂ compositions also resist leakage or transfer of water-soluble components between the aqueous internal phase of the primary emulsion (W₁) and the external aqueous phase (W₂). A W₁-O-W₂ multiple emulsion composition generally is available for immediate application to the skin or hair without the need to shake or agitate the composition in order to redisperse composition ingredients throughout the composition prior to use.

20 In particular, the W₁-O-W₂ multiple emulsion compositions comprise:

- (a) about 40% to about 99% by weight of the composition of an external aqueous phase; and
- (b) about 1% to about 60% by weight of the composition of a primary W₁/O emulsion.

25 The primary W₁/O emulsion comprises:

- (a) about 1% to about 95% by weight of the primary emulsion of an aqueous phase comprising (i) water and (ii) a topically-effective amount of a first topically-active compound, typically about 0.1% to about 30% by weight of the primary emulsion;
- 30 (b) about 0.5% to about 95% by weight of the primary emulsion of an oil phase comprising a volatile silicone compound, a nonvolatile silicone compound, a volatile hydrocarbon compound, a nonvolatile hydrocarbon compound, or a mixture thereof; and
- (c) about 0.1% to about 20% by weight of the primary emulsion of a surfactant phase comprising a silicon-free surfactant or surfactant blend having an HLB value of about 10 or less, and preferably about 7 or less, an oil-soluble silicon-based surfactant, an oil-soluble polymeric surfactant, or a mixture thereof.

35 The external aqueous phase (W₂) is free of a polymeric gelling agent, such as a polyacrylic acid, and optionally includes: (i) 0% to about 15% based on the weight of the external aqueous phase of an emulsifier, such as an anionic surfactant, a cationic surfactant, an amphoteric surfactant, a water-soluble polymeric surfactant, a surfactant system capable of forming stabilizing liquid crystals, a hydrophilic silicon-containing surfactant, a nonionic surfactant or surfactant blend having an HLB value of greater than about 10, or mixtures thereof, and/or (ii) a topically-effective amount of a second topically-active compound, typically 0% to about 30% by weight of the external phase, to perform a function identical to, similar to or different from the first topically-active compound.

40 The W₁-O-W₂ multiple emulsion compositions are stable and exhibit exceptional esthetic and functional properties. The W₁-O-W₂ multiple emulsion compositions are liquids or creams, and are capable of effectively delivering one or more topically-active compounds to the skin or hair from a single composition.

I. The Primary Water-in-Oil (W₁/O) Emulsion

50 The primary W₁/O emulsion comprises water, a first topically-active compound, an oil phase and a surfactant phase. The water and first topically-active compound comprise the aqueous phase of the primary W₁/O emulsion. The primary emulsion comprises droplets containing water and the first topically-active compound (i.e., the aqueous phase). The droplets containing water and the first topically-active compound (W₁) have a diameter ranging from about 1 to about 75 μ and are enveloped by a membrane or film comprising the oil phase and the surfactant phase. The contents of the internal aqueous phase (W₁) therefore do not contact the external aqueous phase (W₂) of the W₁-O-W₂ multiple emulsion.

55 The primary W₁/O emulsion is present in a W₁-O-W₂ multiple emulsion composition in an amount of about 1% to about 60%, and preferably about 5% to about 50%, by weight of the multiple emulsion composition. To achieve the full

advantage of the present invention, the primary W₁/O emulsion is present in an amount of about 10% to about 45%, by weight of the multiple emulsion composition.

A. The Aqueous Phase

5 The aqueous phase is the internal phase (W₁) of the primary W₁/O emulsion. The internal aqueous phase constitutes about 1% to about 95% by weight of the primary emulsion, and the relatively small amounts of oil phase and surfactant phase separate the internal aqueous phase (W₁) of the primary emulsion from the external aqueous phase (W₂).

10 The aqueous phase (W₁) of the present W₁-O-W₂ multiple emulsion compositions comprises water and a first topically-active compound. The aqueous phase can further comprise additional topically-active compounds and/or optional water-soluble compounds capable of providing a desired esthetic or functional effect, such as a fragrance.

15 The aqueous phase (W₁) comprises about 1% to about 95%, and preferably about 10% to about 95%, by weight of the primary emulsion. To achieve the full advantage of the present invention, the aqueous phase comprises about 25% to about 95% by weight of the primary emulsion. Alternatively expressed, the internal aqueous W₁ phase is present in an amount of about 0.01% to about 57%, preferably about 0.5% to about 47.5%, and most preferably about 2.5% to about 42.75%, by weight of the W₁-O-W₂ multiple emulsion composition.

1. Topically-Active Compounds

20 In accordance with an important feature of the present invention, a wide variety of topically-active compounds can be incorporated into the aqueous phase of the primary W₁/O emulsion as the first topically-active compound. The topically-active compounds are water-soluble or water-dispersible and include both cosmetic and medicinal compounds that act upon contact with the skin or hair. The first topically-active compound is present in a sufficient amount to perform its intended function, typically in an amount of about 0.1% to about 30% by weight of the primary emulsion, or, alternatively expressed, about 0.001% to about 17.1% by weight of the W₁-O-W₂ multiple emulsion composition.

25 The first topically-active compound typically remains on the skin or hair after application, as opposed to being rinsed from the skin or hair shortly after application. However, particular first topically-active compounds are designed to be rinsed from the skin or hair in a second rinse step after the compound performs its intended function.

30 In accordance with an important feature of the present invention, the first topically-active compound can be incorporated into the aqueous phase or into the oil phase of the primary emulsion. Whether a particular first topically-active compound is incorporated into the aqueous phase or the oil phase of the primary emulsion is related to the solubility of the topically-active composition in water. In preferred embodiments, the first topically-active compound is water soluble and is incorporated into the internal W₁ aqueous phase.

35 As used herein, the term "water soluble" means water soluble or water dispersible. A water-soluble compound has a water solubility of at least 0.1 g (grams) per 100 ml (milliliters) of water and forms a true solution. A water-soluble compound can be inherently water soluble or can be made water soluble by the addition of a solubilizing compound, such as a coupling agent, a co-surfactant or a solvent. A water-dispersible compound remains dispersed in water for at least the time period necessary to manufacture the primary W₁/O emulsion, i.e., at least about one hour.

40 In addition, the first topically-active compound can be incorporated into the aqueous phase and another topically-active compound can be incorporated into the oil phase to achieve enhanced efficacy (e.g., the primary emulsion includes a medicament in each of the aqueous and oil phases) or to provide different benefits (e.g., the primary emulsion includes a medicament in the aqueous phase and a topical anesthetic in the oil phase).

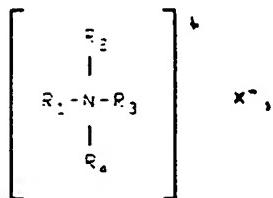
45 The first topically-active compound therefore can be one of, or a combination of, a cosmetic compound, a medicinally-active compound or any other compound that is useful upon topical application to the skin or hair. Such topically-active compounds include, but are not limited to, hair and skin conditioners, hair and skin cleansers, hair fixatives, hair dyes, hair growth promoters, deodorants, skin care compounds, permanent wave compounds, hair relaxers, hair straighteners, antibacterial compounds, antifungal compounds, anti-inflammatory compounds, topical anesthetics, sunscreens and other cosmetic and medicinal topically-effective compounds.

50 In accordance with an important feature of the present invention, the first topically-active compound comprises a water-soluble hair conditioner, such as a quaternary ammonium compound. Quaternary ammonium compounds are substantive to the hair and are excellent hair conditioners, but have a well-known incompatibility with anionic surfactants. Therefore, quaternary ammonium compounds generally are not a component in shampoo-conditioner compositions, but are applied to the hair from a separate conditioning composition after the hair is shampooed.

The water-soluble quaternary ammonium compounds have the general structural formula:

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15 wherein R_1 is an alkyl group including from about 8 to about 18 carbon atoms; R_2 is selected from the group consisting of an alkyl group including from about 8 to about 18 carbon atoms, a hydrogen atom, a methyl group, an ethyl group, a hydroxymethyl group and a hydroxyethyl group; R_3 is selected from the group consisting of a benzyl group, a hydrogen atom, a methyl group, an ethyl group, a hydroxymethyl group and a hydroxyethyl group; R_4 is selected from the group consisting of a hydrogen atom, a methyl group, an ethyl group, a hydroxymethyl group and a hydroxyethyl group; and X^- is an anion. The quaternary nitrogen of the water-soluble quaternary ammonium compound also can be a component of a heterocyclic nitrogen-containing moiety, such as morpholine or pyridine. The anion of the quaternary ammonium compound can be any common anion, such as chloride, methosulfate, ethosulfate, nitrate, bromide, tosylate, acetate, or phosphate.

20 The water-soluble quaternary ammonium compounds have one or two long chain alkyl groups containing from about 8 to about 18 carbon atoms. The long chain alkyl groups also can include, in addition to, or in replacement of, carbon and hydrogen atoms, ether linkages or similar water-solubilizing linkages. The remaining two or three substituents of the quaternary nitrogen of the quaternary ammonium compound can be hydrogen; or benzyl; or short chain alkyl or hydroxyalkyl groups, such as methyl, ethyl, hydroxymethyl or hydroxyethyl groups; or combinations thereof, either of the same or different identity.

25 Exemplary water-soluble quaternary ammonium compounds include, but are not limited to, laurtrimonium chloride; Quaternium-16; lauralkonium chloride; olealkonium chloride; dilauryldimonium chloride; cetalkonium chloride; dicetylmonium chloride; laurylpyridinium chloride; cetylpyridinium chloride; soyatrimonium chloride; Polyquaternium-6; Polyquaternium-7; guarhydroxypropyltrimonium chloride; Polyquaternium-11; Polyquaternium-5; Polyquaternium-10; Polyquaternium-24; cetrimonium chloride; Quaternium-24; mytrimonium chloride; PEG-2 cocomonium chloride; PEG-35 2 cocoyl quaternium-4; PEG-15 cocoyl quaternium-4; PEG-2 stearyl quaternium-4; PEG-15 stearyl quaternium-4; PEG-2 oleyl quaternium-4; and PEG-15 oleyl quaternium-4, and mixtures thereof, wherein the compound designation is provided by the Cosmetic, Toiletry and Fragrance Association, Inc. in the CTFA Cosmetic Ingredient Dictionary, 4th Ed., 1991, hereinafter referred to as the CTFA Dictionary. Other water-soluble quaternary ammonium compounds are listed in the CTFA Cosmetic Ingredient Handbook, 1st Ed., 1988 (hereinafter the CTFA Handbook) at pages 40-42, incorporated herein by reference.

40 Other water-soluble hair conditioners also can be used as the first topically-active compound. Such hair conditioners include, but are not limited to, fatty amine salts, ethoxylated fatty amine salts, dimethicone copolyols, protonated polyethylenimines, protonated ethoxylated polyethylenimines, soluble animal collagen, lauramine oxide, cationic polymers, numerous other water-soluble hair conditioners listed in the CTFA Handbook at pages 71-73, incorporated herein by reference, and mixtures thereof.

45 In addition to hair conditioners, a skin conditioner can be used as the first topically-active compound. Skin conditioning agents include, but are not limited to, humectants, such as fructose, glucose, glycerin, propylene glycol, glycereth-26, mannitol and urea; pyrrolidone carboxylic acid; hydrolyzed lecithin; coco-betaine; cysteine hydrochloride; glutamine; PPG-15; sodium gluconate; potassium aspartate; oleyl betaine; thiamine hydrochloride; sodium laureth sulfate; sodium hyaluronate; hydrolyzed proteins; hydrolyzed keratin; amino acids; amine oxides; water-soluble derivatives of vitamins A, E and D; amino-functional silicones; ethoxylated glycerin; alpha-hydroxy acids and salts thereof; water-soluble fatty oil derivatives, such as PEG-24 hydrogenated lanolin, almond oil, grape seed oil and castor oil; numerous other water-soluble skin conditioners listed in the CTFA Handbook, pages 79-84, incorporated herein by reference; and mixtures thereof.

55 The first topically-active compound also can be a hair fixative or film former that imparts style-retention properties to hair, i.e., sets the hair. The hair fixative typically is a homopolymer, a copolymer, or a terpolymer. The polymers can be nonionic, amphoteric, anionic or cationic. Examples of hair fixatives include, but are not limited to, an acrylamide copolymer; an acrylamide/sodium acrylate copolymer; a polystyrene sulfonate; a polyethylene oxide; a water-dispersible polyester; a cationic cellulose; an acrylate/ammonium methacrylate copolymer; an aminoethylacrylate phosphate/acr-

ylate copolymer; a polyacrylamide; Polyquaternium-1; Polyquaternium-2; Polyquaternium-4; Polyquaternium-5; Polyquaternium-6; Polyquaternium-7; Polyquaternium-8; Polyquaternium-9; Polyquaternium-10; Polyquaternium-11; Polyquaternium-12; Polyquaternium-13; Polyquaternium-14; Polyquaternium-15; Polyquaternium-16; Polyquaternium-28; a PVP (polyvinylpyrrolidone); a PVP/dimethylaminoethylmethacrylate copolymer; a PVP/ethyl methacrylate/methacrylic acid copolymer; a carboxylated polyvinyl acetate; vinyl/caprolactam/PVP/dimethylaminoethyl methacrylate copolymer (GAFFIX VC713); a PVP/vinyl acetate copolymer; a sodium acrylate/vinyl alcohol copolymer; sodium carrageenan; a vinyl acetate/crotonic acid copolymer; numerous other water-soluble hair fixatives listed in the CTFA Handbook at pages 73-74, incorporated herein by reference; and mixtures thereof. Numerous hair fixatives also are disclosed in U.S. Patent No. 5,277,899, incorporated herein by reference.

In addition, the first topically-active compound can be a water-soluble hair dye, such as, but not limited to, m-aminophenol hydrochloride, p-aminophenol sulfate, 2,3-diaminophenol hydrochloride, 1,5-naphthalenediol, p-phenylenediamine hydrochloride, sodium picramate, water-soluble cationic dyes, water-soluble FD&C dyes, like Blue No. 1, Blue No. 2, Red No. 3, Red No. 4, or Red No. 40, water-soluble D&C dyes, like Yellow No. 10, Red No. 22 or Red No. 28, and pyrogallol. Numerous other hair dyes are listed in the CTFA Handbook, pages 70-71, incorporated herein by reference.

The first topically-active compound also can be an antioxidant, like ascorbic acid or erythorbic acid; or a fluorescent whitening agent or optical brightener, like a distyrylbiphenyl derivative, stilbene or a stilbene derivative, a pyralozine derivative or a coumarin derivative. In addition, a self-tanning compound, like dihydroxy acetone, or a hair growth promoter, or a hair bleaching agent, like a perborate or a persulfate salt, can be the first topically-active compound.

The first topically-active compound also can be a deodorant compound, such as an astringent salt or a bioactive compound. The astringent salts include organic and inorganic salts of aluminum, zirconium, zinc, and mixtures thereof. The anion of the astringent salt can be, for example, sulfate, chloride, chlorohydroxide, alum, formate, lactate, benzyl sulfonate or phenyl sulfonate. Exemplary classes of antiperspirant astringent salts include aluminum halides, aluminum hydroxyhalides, zirconyl oxyhalides, zirconyl hydroxyhalides, and mixtures thereof.

Exemplary aluminum salts include aluminum chloride and the aluminum hydroxyhalides having the general formula $\text{Al}_2(\text{OH})_x\text{Q}_y \cdot \text{XH}_2\text{O}$, wherein Q is chlorine, bromine or iodine; x is about 2 to about 5; x+y is about 6, wherein x and y are not necessarily integers; and X is about 1 to about 6. Exemplary zirconium compounds include zirconium oxy salts and zirconium hydroxy salts, also referred to as zirconyl salts and zirconyl hydroxy salts, and represented by the general empirical formula $\text{ZrO}(\text{OH})_{2-nz}\text{L}_z$, wherein z varies from about 0.9 to about 2 and is not necessarily an integer; n is the valence of L; 2-nz is greater than or equal to 0; and L is selected from the group consisting of halides, nitrate, sulfamate, sulfate, and mixtures thereof.

Exemplary deodorant compounds therefore include, but are not limited to, aluminum bromohydrate, potassium alum, sodium aluminum chlorohydroxy lactate, aluminum sulfate, aluminum chlorohydrate, aluminum-zirconium tetrachlorohydrate, an aluminum-zirconium polychlorohydrate complexed with glycine, aluminum-zirconium trichlorohydrate, aluminum-zirconium octachlorohydrate, aluminum sesquichlorohydrate, aluminum sesquichlorohydrex PG, aluminum chlorohydrex PEG, aluminum zirconium octachlorohydrex glycine complex, aluminum zirconium pentachlorohydrex glycine complex, aluminum zirconium tetrachlorohydrex glycine complex, aluminum zirconium trichlorohydrex glycine complex, aluminum chlorohydrex PG, zirconium chlorohydrate, aluminum dichlorohydrate, aluminum dichlorohydrex PEG, aluminum dichlorohydrex PG, aluminum sesquichlorohydrex PG, aluminum chloride, aluminum zirconium pentachlorohydrate, numerous other useful antiperspirant compounds listed in the CTFA Handbook at p. 56, incorporated herein by reference, and mixtures thereof.

In addition to the astringent salts, the deodorant compound can be a bacteriostatic quaternary ammonium compound, such as, for example, cetyl trimethyl ammonium bromide, cetyl pyridinium chloride, benzethonium chloride, diisobutylbenzoxyethoxyethylmethylbenzyl ammonium chloride, sodium N-lauryl sarcosine, sodium N-polymethyl sarcosine, lauroyl sarcosine, N-myristoyl glycine, potassium N-lauroyl sarcosine, and stearyl trimethyl ammonium chloride; or a bioactive compound; or a carbonate or bicarbonate salt, such as, for example, the alkali metal carbonates and bicarbonates, and the ammonium and tetralkylammonium carbonates and bicarbonates.

In addition, other compounds can be included in the primary emulsion as the first topically-active compound in an amount sufficient to perform their intended function. For example, if the composition is intended to be a sunscreen, then compounds such as benzophenone-4, trihydroxycinnamic acid and salts, tannic acid, uric acid, quinine salts, dihydroxy naphtholic acid; an anthranilate, diethanolamine methoxycinnamate, p-aminobenzoic acid, phenylbenzimidazole sulfonic acid, PEG-25 p-aminobenzoic acid or triethanolamine salicylate can be incorporated into the internal aqueous W₁ phase.

Further, sunscreen compounds such as dioxybenzone, ethyl 4-[bis(hydroxypropyl)] aminobenzoate, glyceryl aminobenzoate, homosalate, menthyl anthranilate, octocrylene, octyl methoxycinnamate, octyl salicylate, oxybenzone, padimate O, red petrolatum, titanium dioxide, 4-menthylbenzylidene camphor, benzophenone-1, benzophenone-2, benzophenone-6, benzophenone-12, isopropyl dibenzoyl methane, butyl methoxydibenzoylmethane, zotocrylene, or zinc oxide can be incorporated into the organic phase of the primary emulsion. Other sunscreen compounds soluble in either the aqueous or organic phase are listed in CTFA Handbook, pages 86 and 87, incorporated herein by reference.

Similarly, topically-active drugs, like antifungal compounds; antibacterial compounds; anti-inflammatory compounds; topical anesthetics; skin rash, skin disease and dermatitis medications; and anti-itch and irritation-reducing compounds can be included in the compositions of the present invention. For example, analgesics such as benzocaine, dyclonine hydrochloride, aloe vera and the like; anesthetics such as butamben picrate, lidocaine hydrochloride, xylocaine and the like; antibacterials and antiseptics, such as povidone-iodine, polymyxin b sulfate-bacitracin, zinc-neomycin sulfate-hydrocortisone, chloramphenicol, methylbenzethonium chloride, and erythromycin and the like; antiparasitics, such as lindane; deodorants, such as chlorophyllin copper complex, aluminum chloride, aluminum chloride hexahydrate, and methylbenzethonium chloride; essentially all dermatologicals, like acne preparations, such as benzoyl peroxide, erythromycin-benzoyl peroxide, clindamycin phosphate, 5,7-dichloro-8-hydroxyquinoline, and the like; anti-inflammatory agents, such as alclometasone dipropionate, betamethasone valerate, and the like; burn relief ointments, such as o-amino-p-toluenesulfonamide monoacetate and the like; depigmenting agents, such as monobenzene; dermatitis relief agents, such as the active steroid amcinonide, diflorasone diacetate, hydrocortisone, and the like; diaper rash relief agents, such as methylbenzethonium chloride and the like; emollients and moisturizers, such as mineral oil, PEG-4 dilaurate, lanolin oil, petrolatum, mineral wax and the like; fungicides, such as butocouazole nitrate, haloprogin, clotrimazole, and the like; herpes treatment drugs, such as O-[(2-hydroxyethoxy)methyl]guanine; pruritic medications, such as alclometasone dipropionate, betamethasone valerate, isopropyl myristate MSD, and the like; psoriasis, seborrhea and scabicide agents, such as anthralin, methoxsalen, coal tar and the like; steroids, such as 2-(acetoxy)-9-fluoro-1',2',3',4'-tetrahydro-11-hydroxypregna-1,4-dieno[16,17-b]naphthalene-3,20-dione and 21-chloro-9-fluoro-1',2',3',4'-tetrahydro-11b-hydroxypregna-1,4-dieno[16z,17-b]naphthalene-3,20-dione. Any other medication capable of topical administration also can be incorporated in a composition of the present invention in an amount sufficient to perform its intended function. Other topically-active compounds are listed in Remington's Pharmaceutical Sciences, 17th Ed., Merck Publishing Co., Easton, PA (1985), pages 773-791 and pages 1054-1058 (hereinafter Remington's), incorporated herein by reference.

An above-described first topically-active compound is designed to remain on the skin or hair to perform its intended function. However, in particular situations, a first topically-active compound that is rinsed from the skin or hair can be incorporated into the internal aqueous W₁ phase of the primary emulsion.

For example, a W₁-O-W₂ multiple emulsion composition designed as a permanent wave composition can incorporate a reducing agent into the external aqueous phase as the second topically-active compound. After applying the W₁-O-W₂ composition to the hair and allowing the composition to contact the hair for a sufficient time to reduce the hair, the external aqueous phase is rinsed from the hair leaving droplets of the primary emulsion on the hair.

The primary emulsion has incorporated therein an oxidizing agent as the first topically-active compound. After the oil phase, and preferably a volatile oil phase, evaporates from the primary emulsion, the oxidizing agent is released to neutralize the hair and any reducing agent remaining on the hair. Excess oxidizing agent then can be rinsed from the hair in a second rinsing step.

Exemplary, but non-limiting, oxidizing agents used as the first topically-active compound are ammonium persulfate, hydrogen peroxide, potassium bromate, potassium chromate, potassium persulfate, sodium bromate, sodium carbonate peroxide, sodium iodate, sodium perborate, sodium persulfate, urea peroxide, and mixtures thereof. An oxidizing agent also can be the second topically-active compound when the first topically-active compound is a hair conditioner. In addition, the second topically-active compound can be a bleaching agent (i.e., an oxidizing agent) and the first topically-active compound can be a hair dye.

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2. Water

Sufficient water is present in the aqueous phase such that the aqueous phase comprises about 1% to about 95% by weight of the primary emulsion. Total water present in the W₁-O-W₂ multiple emulsion composition is about 30% to about 99.9%, and typically about 40% to about 95%, by weight of the composition.

3. Optional Ingredients

The internal aqueous phase also can include optional ingredients traditionally included in topically-applied compositions. These optional ingredients include, but are not limited to, dyes, fragrances, preservatives, antioxidants, detackifying agents, and similar types of compounds. The optional ingredients are included in the internal aqueous phase of the primary emulsion in an amount sufficient to perform their intended function.

B. The Oil Phase

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The primary W₁/O emulsion also comprises about 0.5% to about 95%, preferably about 0.5% to about 80%, and most preferably about 0.5% to about 75%, by weight of the primary emulsion of an oil phase; alternatively expressed as about 0.005% to about 57%, preferably about 0.005% to about 48%, and most preferably about 0.005% to about 45%, by weight of the W₁-O-W₂ multiple emulsion composition. The oil phase encapsulates the internal aqueous W₁

phase of the primary emulsion to form droplets of about 5 to about 1,000 μ , and preferably about 10 to about 500 μ , in diameter. The oil phase therefore provides a barrier between the internal aqueous phase of the primary emulsion and the external aqueous phase of the W₁-O-W₂ multiple emulsion composition. The oil phase provides a sufficient barrier even though the oil phase constitutes as little as 0.5% by weight of the primary emulsion.

5 The oil phase can be a volatile oil phase, a nonvolatile oil phase, or a mixture thereof. A volatile oil phase comprising a volatile silicone compound, a volatile hydrocarbon compound, or a mixture thereof is preferred, and evaporates from the skin or hair after rinsing to release the first topically-active compound. A nonvolatile oil phase, comprising a nonvolatile silicone compound, a nonvolatile hydrocarbon compound, or a mixture thereof, remains on the skin or hair after the rinsing step and the first topically-active compound is released by rubbing the skin or hair to break the primary W₁/O emulsion.

10 In accordance with an important feature of the present invention, the oil phase comprises a combination of a volatile oil phase and a nonvolatile oil phase. In this embodiment, an oil phase can be designed to evaporate at a preselected temperature and provide a controlled release of the first topically-active compound at the preselected temperature. Preselected temperatures are those encountered during normal hair drying, provided by a hair dryer, or provided by a
15 curling iron.

15 In one embodiment, the oil phase is sufficiently volatile to evaporate during the process of drying the skin or hair, and thereby release the internal aqueous W₁ phase, which includes the first topically-active compound, to contact the skin or hair. The volatile oil phase evaporates after the external aqueous phase of the W₁-O-W₂ multiple emulsion composition is rinsed from the skin or hair.

20 As previously stated, the oil phase also can include a water-insoluble topically-active compound in a sufficient amount to impart a particular functional or esthetic effect (e.g., emolliency), as long as the topically-active compound does not adversely affect the W₁-O-W₂ multiple emulsion composition (e.g., does not impart emulsion instability). Although the oil phase can incorporate a topically-active compound, the topically-active compound preferably is incorporated into the internal aqueous phase.

25 The oil phase can comprise a volatile oil phase, a nonvolatile oil phase or a mixture thereof. The volatile oil phase comprises a volatile silicone compound, a volatile hydrocarbon compound or a mixture thereof. Preferably, the volatile oil phase comprises a volatile silicone compound. The volatile oil phase evaporates from the skin or hair to release the first topically-active compound.

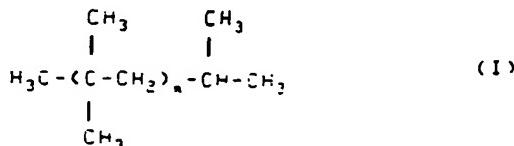
Exemplary volatile compounds include, but are not limited to, volatile, low molecular weight polydimethylsiloxane
30 compounds. The volatile, low molecular weight polydimethylsiloxane compound can be either a linear or a cyclic polydimethylsiloxane compound, as long as the polydimethylsiloxane compound has sufficient volatility to volatilize after topical application to the skin or hair. Preferably the polydimethylsiloxane is a cyclic siloxane, like cyclomethicone. The volatile silicone evaporates relatively quickly to leave only the nonvolatile components of the primary emulsion on the skin. In general, volatile polydimethylsiloxane compounds useful in the compositions of the present invention include
35 polydimethylsiloxane compounds having a viscosity of about 0.5 to about 10 cs (centistokes). The preferred volatile polydimethylsiloxanes have a viscosity in the range of about 0.5 to about 6 cs.

The cyclic, low molecular weight, volatile polydimethylsiloxanes, designated in the CTFA International Cosmetic Ingredient Dictionary, 4th Ed., Cosmetic, Toiletry and Fragrance Association, Washington, DC (1991) (hereinafter CTFA Dictionary) as cyclomethicones, are the preferred siloxanes used in a composition of the present invention. To achieve
40 the full advantage of the present invention, a cyclomethicone used in a composition of the present invention is a low viscosity, low molecular weight, water-insoluble cyclic compound having an average of about 3 to about 6 -[O-Si(CH₃)₂]-repeating group units per molecule (hexamethylcyclotrisiloxane, octamethylcyclotetrasiloxane, decamethylcyclopentasiloxane, dodecamethylcyclohexasiloxane, and mixtures thereof); boil at atmospheric pressure at about 150°C to about 250°C; and have a viscosity at 25°C of about 2 to about 6 centistokes. The polydimethyl cyclosiloxanes having an average
45 of about 4 to about 5 repeating units per molecule, i.e., the tetramer and pentamer, are especially preferred. To achieve the full advantage of the present invention, the volatile cyclomethicone has a boiling point at atmospheric pressure in the range of about 150°C to about 220°C. Suitable cyclomethicones are available commercially under the tradenames DOW CORNING 245 FLUID, DOW CORNING 344 FLUID and DOW CORNING 345 FLUID from Dow Corning Corporation, Midland, Mich., and SILICONE SF-1173 and SILICONE SF-1202 from General Electric, Waterford, N.Y.

50 An example of a linear, low molecular weight, volatile polydimethylsiloxane compound useful in the composition and method of the present invention is the compound designated in the CTFA Dictionary as hexamethydisiloxane, available commercially under the tradename DOW CORNING 200 FLUID from Dow Corning Corp., Midland, Mich. Hexamethyl-disiloxane has a viscosity of 0.65 cs and is highly volatile. Other linear polydimethylsiloxanes, such as decamethyltetrasiloxane, having a boiling point of about 195° and a viscosity of 1.5 cs; octamethyltrisiloxane; and dodecamethylpentasiloxane, also have sufficient volatility to be useful in the composition of the present invention. Another useful linear siloxane is bisphenylhexamethicone. In general, it has been found that a linear, low molecular weight volatile polydimethylsiloxane compound having a viscosity at 25°C and atmospheric pressure of about 0.5 to about 5 cs, and a boiling point at atmospheric pressure of about 100°C to about 250°C, is preferred for use in the composition and method of the present invention.

The volatile oil phase also can comprise a volatile hydrocarbon compound, such as a hydrocarbon having about 10 to about 30 carbon atoms, that has sufficient volatility to slowly volatilize from the skin or hair after application of the W₁-O-W₂ multiple emulsion composition to the skin or hair and subsequent rinsing. A preferred volatile hydrocarbon compound is an aliphatic hydrocarbon having about 12 to about 24 carbon atoms, and having a boiling point of about 100°C to about 250°C. The volatile hydrocarbon compounds perform the same function and provide essentially the same benefits as the volatile silicone compounds.

Volatile hydrocarbon compounds incorporated into the primary emulsion include, for example, isododecane and isohexadecane, i.e., PERMETHYL 99A, PERMETHYL 101A and PERMETHYL 102A, available from Presperse, Inc., South Plainfield, NJ. Other exemplary volatile hydrocarbon compounds are depicted in general structural formula (I), wherein n ranges from 2 to 5.



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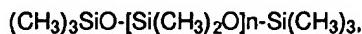
Another exemplary volatile hydrocarbon compound is ISOPAR M (a C₁₃-C₁₄ isoparaffin available from Exxon Chemical Co., Baytown, TX).

As previously stated, the oil phase also can be a nonvolatile oil phase. The nonvolatile oil phase comprises a non-volatile silicone compound, a nonvolatile hydrocarbon, or a mixture thereof. Preferably, the nonvolatile oil phase comprises a nonvolatile silicone compound. The nonvolatile oil phase does not evaporate from the skin or hair. The first topically-active compound therefore is released by rubbing the skin or hair to rupture the primary W₁/O emulsion. The nonvolatile oil phase boils at atmospheric pressure at a temperature above about 250°C.

Exemplary nonvolatile silicone compounds include a polyalkyl siloxane, a polyaryl siloxane or a polyalkylaryl siloxane. Mixtures of these nonvolatile silicone compounds also are useful. The nonvolatile silicones are nonfunctional siloxanes or siloxane mixtures having a viscosity of about 10 to about 600,000 cs, and typically about 350 to about 10,000 cs, at 25°C. The so-called "rigid silicones", as described in U.S. Patent No. 4,902,499, herein incorporated by reference, having a viscosity above 600,000 cs at 20°C, and a weight average molecular weight of at least about 500,000, also are useful in a composition of the present invention. Phenyltrimethicone also can be used as the nonvolatile silicone compound.

The preferred nonvolatile silicone compound is a nonvolatile polydimethylsiloxane compound, such as a mixture, in about a 2:1 weight ratio, of a low molecular weight polydimethylsiloxane fluid and a higher molecular weight polydimethylsiloxane gum. Preferred silicone gums include linear and branched polydimethylsiloxanes of the following general formula:

40



wherein n is a number from about 2,000 to about 15,000, and preferably from about 2,000 to about 7,000. Silicone gums useful in compositions of the present invention are available from a variety of commercial sources, including General Electric Company, Waterford, NY and Dow Corning Corp., Midland, MI.

The nonvolatile oil phase also can comprise a nonvolatile hydrocarbon compound, such as mineral oil. Other exemplary nonvolatile hydrocarbon compounds that can be incorporated into the oil phase include, but are not limited to, a branched 1-decene oligomer, like 1-decene dimer or a polydecene.

The oil phase also optionally can comprise (1) an oil, such as jojoba oil, wheat germ oil or purcellin oil; or (2) a water-insoluble emollient, such as, for example, an ester having at least about 10 carbon atoms, and preferably about 10 to about 32 carbon atoms.

Suitable esters include those comprising an aliphatic alcohol having about eight to about twenty carbon atoms and an aliphatic or aromatic carboxylic acid including from two to about twelve carbon atoms, or conversely, an aliphatic alcohol having two to about twelve carbon atoms with an aliphatic or aromatic carboxylic acid including about eight to about twenty carbon atoms. The ester is either straight-chained or branched. Preferably, the ester has a molecular weight of less than about 500. Suitable esters therefore include, for example, but are not limited to:

- (a) aliphatic monohydric alcohol esters, including but not limited to:
myristyl propionate,

- isopropyl isostearate,
 isopropyl myristate,
 isopropyl palmitate,
 5 cetyl acetate,
 cetyl propionate,
 cetyl stearate,
 isodecyl neopentanoate,
 cetyl octanoate,
 isocetyl stearate;
- 10 (b) aliphatic di- and tri-esters of polycarboxylic acids, including but not limited to:
 diisopropyl adipate,
 diisostearyl fumarate,
 dioctyl adipate, and
 triisostearyl citrate;
- 15 (c) aliphatic polyhydric alcohol esters, including but not limited to:
 propylene glycol dipelargonate;
- (d) aliphatic esters of aromatic acids, including but not limited to:
 C₁₂-C₁₅ alcohol esters of benzoic acid,
 20 octyl salicylate,
 sucrose benzoate, and
 dioctyl phthalate.

Numerous other esters are listed in the CTFA Handbook, at pages 24 through 26, incorporated herein by reference.

25 **C. The Surfactant Phase**

The primary emulsion of the present invention also includes about 0.1% to about 20%, and preferably about 0.1% to about 15%, by weight of the primary emulsion of a surfactant phase. To achieve the full advantage of the present invention, about 0.5% to about 12% by weight of the surfactant phase is present in the primary emulsion. Alternatively expressed, the surfactant phase is present in an amount of about 0.001% to about 12%, preferably about 0.005% to about 7.5%, and most preferably about 0.05% to about 5.4%, by weight of the W₁-O-W₂ multiple emulsion composition.

The surfactant phase comprises a silicon-free surfactant, or a blend of silicon-free surfactants, having an HLB value of about 10 or less (i.e., an HLB value of about 0.1 to about 10), an oil-soluble silicon-based surfactant, an oil-soluble polymeric surfactant, or mixtures thereof. Preferably, the silicon-free surfactant or surfactant blend has an HLB value of about 1 to about 7. To achieve the full advantage of the present invention, the silicon-free surfactant or surfactant blend has an HLB value of about 3 to about 6. The term "oil-soluble" as used herein means a compound having a solubility of at least 0.1 g per 100 ml of oil phase to form a true solution.

The surfactant phase therefore comprises a single silicon-free surfactant having an HLB value of about 0.1 to about 10, or a blend of silicon-free surfactants having different HLB values such that the blend has an HLB value of about 1 to about 10. The surfactant phase also can comprise a blend of an oil-soluble silicon-based surfactant, or an oil-soluble polymeric surfactant, and a silicon-free surfactant or surfactant blend having an HLB value of about 10 or less. The above-described surfactant phases provide a water-in-oil emulsion.

Preferably, the surfactant phase comprises an oil-soluble silicon-based surfactant when the oil phase comprises a volatile or a nonvolatile silicone compound. If the oil phase is a volatile or nonvolatile hydrocarbon compound, the surfactant phase preferably comprises a silicon-free nonionic surfactant or an oil-soluble polymeric surfactant. If a combination of a silicone compound and hydrocarbon compound is used as the oil phase, the surfactant phase preferably comprises a combination of (1) a silicon-based surfactant and (2) an oil-soluble silicon-free surfactant, an oil-soluble polymeric surfactant or a mixture thereof.

A silicon-free nonionic surfactant having an HLB value of about 0.1 to about 10 can be used alone as the surfactant phase of the present invention. The surfactant phase also can comprise a blend of silicon-free surfactants each having an HLB value of less than 10. In addition, silicon-free surfactants having an HLB value of about 0.1 to about 10 also can be used as the first surfactant of a surfactant blend having an HLB value of about 1 to about 10, then a silicon-free surfactant having an HLB of greater than about 10 is the second surfactant of the surfactant blend having an HLB of about 1 to about 10.

55 Typically, silicon-free nonionic surfactants having an HLB value of about 10 or less have a hydrophobic moiety, such as a long chain (C₈-C₂₂) alkyl group or an alkylated aryl group, and a hydrophilic chain comprising a small number (i.e., one to about six) of ethoxy moieties or a combination of ethoxy and propoxy moieties. The silicon-free nonionic surfactants having an HLB of greater than about 10 typically have the same type of hydrophobic moiety as the low HLB surfactants, but include more ethoxy and/or propoxy moieties.

The HLB value of a particular silicon-free surfactant can be found in McCutcheon's Emulsifiers and Detergents, North American and International Editions, MC Publishing Co., Glen Rock, NJ (1993) (hereinafter McCutcheon's). Alternatively, the HLB value of a particular surfactant can be estimated by dividing the weight percent of oxyethylene in the surfactant by five (for surfactants including only ethoxy moieties). In addition, the HLB value of a surfactant blend can be estimated by the following formula:

$$\text{HLB} = (\text{wt.\%A}) (\text{HLB}_A) + (\text{wt.\%B}) (\text{HLB}_B),$$

wherein wt.% A and wt.% B are the weight percent of surfactants A and B in the silicon-free surfactant blend, and HLB_A and HLB_B are the HLB values for surfactants A and B, respectively.

Exemplary classes of silicon-free nonionic surfactants include, but are not limited to, polyoxyethylene ethers of fatty (C_6 - C_{22}) alcohols, polyoxyethylene/polyoxypropylene ethers of fatty (C_6 - C_{22}) alcohols, ethoxylated alkylphenols, polyethylene glycol ethers of methyl glucose, polyethylene glycol ethers of sorbitol, and mixtures thereof.

Exemplary silicon-free nonionic surfactants are the ethoxylated alcohols having an HLB value of about 0.1 to about 10. An especially preferred ethoxylated alcohol is laureth-1, i.e., lauryl alcohol ethoxylated with an average of one mole of ethylene oxide. Other suitable ethoxylated alcohols include laureth-2, laureth-3 and laureth-4. Numerous other non-ionic surfactants having an HLB of about 0.1 to about 10 are listed in McCutcheon's at pages 229-236, incorporated herein by reference. Other exemplary silicon-free nonionic surfactants having an HLB value of about 0.1 to about 10 include, but are not limited to, the ethoxylated nonylphenols, ethoxylated octylphenols, ethoxylated dodecylphenols, ethoxylated fatty (C_6 - C_{22}) alcohols having four or fewer ethylene oxide moieties, oleth-2, steareth-3, steareth-2, ceteth-2, oleth-3, and mixtures thereof.

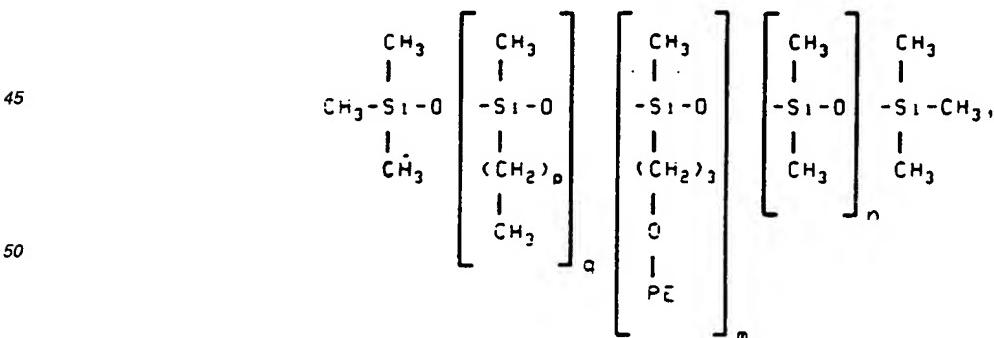
The surfactant phase also can comprise a silicon-free surfactant blend having an HLB value of about 1 to about 10. The blend is a mixture of a sufficient amount of a surfactant having a low HLB value, i.e., about 0.1 to about 10, and a sufficient amount of a surfactant having a higher HLB value, i.e., about 1 to greater than about 10, such that the surfactant blend has an HLB value of about 1 to about 10. Exemplary, but nonlimiting, nonionic surfactants having a high HLB value are listed in McCutcheon's at pages 236-246, incorporated herein by reference.

Exemplary silicon-free nonionic surfactants having an HLB value greater than about 10 are oleth-10, octylphenol or nonylphenol ethoxylated with six or more moles of ethylene oxide, steareth-10, trideceth-6, methyl gluceth-10, dodoxynol-12, ceteth-12, C_{11-15} pareth-20, and mixtures thereof. The identity of the high HLB surfactant is not limited as long as the resulting surfactant phase provides a stable water-in-oil primary emulsion.

An exemplary oil-soluble silicon-based surfactant is a dimethicone copolyol, which is a dimethylsiloxane polymer having polyoxyethylene and/or polyoxypropylene side chains, such as DOW CORNING 3225C FORMULATION AID, available from Dow Corning Co., Midland, MI. The dimethicone copolyol has about 15 or fewer ethylene oxide and/or propylene oxide monomer units, in total, in the side chains. Dimethicone copolyols conventionally are used in conjunction with silicones because the silicon-containing surfactants are extremely soluble in a volatile or a nonvolatile silicone compound, are extremely insoluble in water, and have a low skin irritancy potential.

Another exemplary, but nonlimiting, oil-soluble, silicon-based surfactant is an alkyl dimethicone copolyol, such as cetyl dimethicone copolyol available commercially as ABIL® EM 90 from Goldschmidt Chemical Corporation, Hopewell, VA. The alkyl dimethicone copolyols have the structure:

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wherein p is a numeral from 7 through 17;

q is a numeral from 1 through 100;

m is a numeral from 1 through 40;

n is a numeral from 0 through 200; and

PE is $(C_2H_4O)_a(C_3H_6O)_b-H$ having a molecular weight of about 250 to about 2000, wherein a and b are selected such that the weight ratio of C_2H_4O/C_3H_6O is from 100/0 to 20/80.

The surfactant phase also can comprise an oil-soluble polymeric surfactant. Polymeric surfactants capable of forming water-in-oil emulsions completely cover the surface of the water droplet, are firmly anchored at the oil-water interface, the external oil phase is a good solvent for the stabilizing portion of the polymeric surfactant and the thickness of the polymer layer on the oil side of the interface is sufficient to ensure stability. These surfactants may include ethoxy, propoxy and/or similar alkylene oxide monomer units, e.g., butoxy. The oil-soluble polymeric surfactants act as surfactants and are not physically or chemically crosslinked in solution. The oil-soluble polymeric surfactants therefore are differentiated from polymeric gelling agents such as polyacrylic acid or polymethacrylic acid.

Accordingly, exemplary oil-soluble polymeric surfactants include, but are not limited to, polyoxyethylene-polyoxypropylene block copolymers, and similar polyoxyalkylene block copolymers. The oil-soluble block copolymers typically have less than about 20% by weight of ethylene oxide. Specific nonlimiting oil-soluble polymeric surfactants include Poloxamer 101, Poloxamer 105, PPG-2-Buteth-3, PPG-3-Buteth-5, PPG-5-Buteth-7, PPG-7-Buteth-10, PPG-9-Buteth-12, PPG-12-Buteth-16, PPG-15-Buteth-20, PPG-20-Buteth-30, PPG-24-Buteth-27, PPG-28-Buteth-35, and PEG-15 Butanediol. Other useful oil-soluble polymeric surfactants are poloxamines, i.e., polyoxyethylene-polyoxypropylene block copolymers of ethylene diamine, having less than about 40% by weight ethylene oxide.

In accordance with an important feature of the present invention, the hydrophobic moiety of a silicon-free surfactant, silicon-containing surfactant or polymeric surfactant is sufficiently soluble in the oil phase such that a sufficient amount of the surfactant is present in the oil phase to stabilize the primary W₁/O emulsion. In one embodiment, when the oil phase comprises a silicone compound, the surfactant phase comprises either a silicon-based surfactant, a silicon-free surfactant having a hydrophobic moiety preferably containing about ten to about fourteen carbon atoms, an oil-soluble polymeric surfactant, or a mixture thereof. If the hydrophobic moiety of the silicon-free surfactant is saturated and includes more than about 14 carbon atoms, the silicon-free surfactant is insoluble in the silicone phase and the primary W₁/O emulsion is unstable. If the hydrophobic moiety includes less than about 10 carbon atoms, the primary W₁/O emulsion has a tendency to coalesce, i.e., the emulsion droplets fuse to form large droplets. The amount of surfactant phase necessary to provide a primary emulsion of desired W₁/O droplet diameter varies with the amount of aqueous phase in the primary emulsion and is easily determined by those skilled in the art.

30 II. The External Aqueous Phase

The external aqueous phase (W₂) of the multiple emulsion can comprise only water. In this embodiment, the primary W₁/O emulsion is stable, but the W₁-O-W₂ multiple emulsion composition is not phase stable, separating into two phases during storage (i.e., separates into the external aqueous phase and the primary emulsion). In this embodiment, the W₁-O-W₂ multiple emulsion composition requires shaking to redisperse the primary emulsion in the continuous external aqueous phase prior to applying the composition to the skin or hair.

In addition, in this embodiment, the continuous external aqueous phase provides no cosmetic or medicinal benefits to the skin or hair. If necessary, the water-only W₂ external phase is rinsed from the hair. Irrespective of a rinsing step, the treated hair is dried to evaporate the rinse water or external aqueous phase and the volatile oil phase, and thereby release the first topically-active compound in the internal aqueous phase of the primary emulsion to contact the skin or hair.

Such a composition typically has a low consumer acceptance because the presence of two distinct phases is esthetically unappealing and because the composition must be shaken prior to each use. However, such a composition provides the advantages of favorable economics and an ability to apply a water-soluble topically-active compound to the skin or hair from a rinse-off product. The large amount of water-soluble, topically-active compound typically wasted in the rinsing step is reduced substantially, which permits formulators to incorporate less of the topically-active compound into the rinse-off composition.

In a preferred embodiment, the external aqueous phase further comprises an emulsifier in an amount of 0% to about 15%, and preferably about 0.1% to about 10%, by weight of the external aqueous phase; alternatively expressed as 0% to about 14.85%, preferably about 0.001% to about 9.9%, by weight of the W₁-O-W₂ multiple emulsion composition. The emulsifier is present in a sufficient amount to provide a stable W₁-O-W₂ multiple emulsion composition, wherein droplets of the primary W₁/O emulsion are uniformly distributed throughout the external aqueous phase, and preferably remain uniformly distributed in the external aqueous phase throughout the life of the multiple emulsion composition. A sufficient amount of surfactant in the external aqueous phase provides W₁/O droplets of a sufficiently large size (i.e., about 0.5 to about 1000 μ) to deposit on the skin or hair, and to resist rinsing from the skin or hair.

A W₁-O-W₂ multiple emulsion composition is tested for stability immediately after the W₁-O-W₂ multiple emulsion composition is prepared. The composition is observed under an optical microscope, and the W₁-O-W₂ multiple emulsion composition is defined as stable when droplets of the primary W₁/O emulsion are observed at a magnification of 50 or

greater. The W₁-O-W₂ multiple emulsion composition then is examined at periodic intervals for extended stability. Preferably, the W₁-O-W₂ multiple emulsion composition is stable for the expected product life of the skin or hair care product.

The emulsifier comprises an anionic surfactant, a cationic surfactant, an amphoteric surfactant, a water-soluble polymeric surfactant, a water-soluble silicon-containing surfactant, a nonionic surfactant having an HLB of greater than about 10, or a surfactant system capable of forming stabilizing liquid crystals around the W₁/O droplets. The nonionic surfactant preferably has an HLB of at least about 12, and more preferably, an HLB value of at least about 15.

The emulsifier for the external phase does not gel the external aqueous phase. The emulsifier however may be capable of forming a stabilizing layer of lamellar liquid crystals around droplets of the internal primary W₁/O emulsion. This barrier film prevents coalescence between primary emulsion droplets and precludes migration of droplet contents from the primary emulsion to the external aqueous phase.

The emulsifier included in the external aqueous phase has a low affinity for the oil phase. The emulsifier therefore does not disrupt the primary W₁/O emulsion and avoids the formation of a simple oil-in-water emulsion. Accordingly, the emulsifier in the external aqueous phase comprises a nonionic surfactant or surfactant blend having an HLB value of greater than about 10, a cationic surfactant, an anionic surfactant, an amphoteric surfactant, a water-soluble polymeric surfactant, a water-soluble silicon-containing surfactant, a surfactant system capable of forming stabilizing liquid crystals, or a compatible mixture thereof.

Exemplary nonionic surfactants having an HLB value of greater than about 10 include, but are not limited to, trideceth-6, ceteth-10, laureth-9, octoxynol-9, nonoxynol-12, a poloxamer, trideceth-12, oleth-20 and polysorbate-20. Numerous other nonionic surfactants having an HLB value of greater than about 10 are listed in McCUTCHEON'S, pages 235-246, incorporated herein by reference.

Exemplary cationic surfactants include, but are not limited to, lauryl dimethyl amine oxide, stearyl dimethyl amine oxide, coco amidopropyl dimethylamine oxide, stearamidopropyl dimethylamine, C₁₄₋₂₀ isoalkylamidopropylethyldimmonium ethosulfate, C₁₈₋₂₂ isoalkylamidopropylethyldimonium ethosulfate, cocamidopropylmonium hydroxypropylamino hydrolyzed animal protein, hydroxyanthraquinoneaminopropyl methyl morpholinium methosulfate, oleamidopropyl ethyldimonium ethosulfate, rapeseedamidopropyl benzylmonium chloride, rapeseedamidopropyl ethyldimonium ethosulfate, ricinoleamidopropyl ethyldimonium ethosulfate, soyamidopropyl benzylmonium chloride, soyamidopropyl ethyldimonium ethosulfate, stearamidopropalkonium chloride, stearamidopropyl cetearyl dimonium tosylate, and mixtures thereof. Also, the quaternary ammonium compounds, protonated polyethyleneimines, protonated fatty amines and other cationic hair conditioners disclosed as the first topically-active compound can be used as the emulsifier in the external aqueous phase.

Exemplary anionic surfactants include, but are not limited to, compounds having a hydrophobic moiety, such as a carbon chain including from about eight carbon atoms to about 30 carbon atoms, and particularly from about twelve carbon atoms to about twenty carbon atoms; and further includes a hydrophilic moiety, such as sulfate, sulfonate, carbonate, phosphate or carboxylate. Often, the hydrophobic carbon chain is etherified, such as with ethylene oxide or propylene oxide.

Suitable anionic surfactants include compounds in the classes known as alkyl sulfates, alkyl ether sulfates, alkylamide sulfates, alkylamide ether sulfates, alkyl ether sulfonates, alkylamide sulfonates, sulfate esters of an alkylphenoxy polyoxyethylene ethanol, alpha-olefin sulfonates, beta-alkyloxy alkane sulfonates, alkyl arylsulfonates, alkyl carbonates, alkyl ether carboxylates, fatty acids, soaps, alkyl sulfosuccinates, alkyl ether sulfosuccinates, alkylamide sulfosuccinates, sarcosinates, alkyl phosphates, octoxynol or nonoxynol phosphates, taurates, fatty taurides, sulfated monoglycerides, fatty acid amido polyoxyethylene sulfates, isethionates, and mixtures thereof.

Usually, the anionic surfactant is present in the composition as a neutralized salt in the form of a sodium, potassium, lithium, ammonium, alkylammonium or hydroxyalkylammonium salt, wherein the alkyl moiety includes from one to about three carbon atoms. The alkyl sulfates and alkyl ether sulfates are particularly effective classes of anionic surfactants. Consequently, exemplary anionic surfactants include, but are not limited to, the ammonium, monoethanolamine, diethanolamine, triethanolamine, isopropylamine, sodium, potassium, lithium or magnesium salt of lauryl sulfate, dodecylbenzenesulfonate, lauryl sulfosuccinate, lauryl ether sulfate, lauryl ether carboxylate, lauryl sarcosinate, cocomethyl tauride, sulfosuccinate half ester amide, and mixtures thereof. Examples of especially useful anionic surfactants are a lauryl sulfate salt and a lauryl ether sulfate salt.

Exemplary amphoteric surfactants include, but are not limited to, cocoamidopropyl dimethyl betaine, cocoamido-propyl hydroxysultaine, cocodimethyl betaine, coco imidazoline dicarboxylate, coco imidazoline monocarboxylate, coco-betaine, lauryl chloride, lauryl sultaine, decyl betaine, oleamidopropyl betaine, tallowamidopropyl betaine, sodium caproamphacetate, sodium lauroamphacetate, sodium cocoamphopropylsulfonate, numerous other alkylamido alkylamines and betaines listed in the CTFA Handbook, at pages 9, 10, 15 and 16, incorporated herein by reference, and mixtures thereof.

Many additional nonionic, cationic, anionic, and amphoteric surfactants are listed in McCUTCHEON'S EMULSIFIERS AND DETERGENTS, 1993 ANNUAL, published by McCutcheon Division, MC Publication Co., Glen Rock, NJ, and in the CTFA Handbook, each incorporated herein by reference.

The water-soluble polymeric surfactants useful in the external aqueous phase (W_2) are different from the oil-soluble polymeric surfactants useful in the surfactant phase of the primary W_1/O emulsion. The water-soluble polymeric surfactants useful in the external aqueous phase (W_2) stabilize the W_1-O-W_2 multiple emulsion composition to maintain composition stability and reduce phase separation between the dispersed primary W_1/O emulsion and the continuous external aqueous phase (W_2). The water-soluble polymeric surfactants do not gel the external phase of the W_1-O-W_2 multiple emulsion composition because the surfactants do not physically or chemically crosslink in solution.

In particular, a water-soluble polymeric surfactant useful in the external aqueous phase (W_2) of the multiple emulsion composition completely covers the surface of the oil droplet, is firmly anchored at the oil-water interface, the external water phase is a good solvent for the stabilizing portion of the polymeric surfactant, and the thickness of the polymer layer on the water side of the interface is sufficient to ensure stability. These surfactants can include repeating ethoxy, propoxy and/or similar alkylene oxide monomer units. Exemplary water-soluble polymeric surfactants include, but are not limited to, polyoxyethylene-polyoxypropylene block copolymers, and similar polyoxyalkylene block copolymers. The water-soluble block copolymers typically include 20%, by weight, or more of ethylene oxide. Specific nonlimiting water-soluble polymeric surfactants include Poloxamer 108 through Poloxamer 407 and PPG-33-Buteth-45. Other useful water-soluble polymeric surfactants are poloxamines having at least 40%, by weight, or more ethylene oxide.

The water-soluble silicon-containing surfactants useful in the external phase (W_2) are different from the oil-soluble silicon-based surfactants used in the surfactant phase of the primary W_1/O emulsion. The water-soluble silicon-containing surfactants are dimethicone copolyols having polyoxyethylene and/or polyoxypropylene side chains having about 15 or more total ethylene oxide and propylene oxide monomer units.

In addition, a surfactant system capable of forming a liquid crystal structure also can be used as the emulsifier in the external aqueous phase (W_2). The surfactant system can be a single surfactant, or a blend of surfactants. In some cases, a particular surfactant cannot form a liquid crystal structure alone, but can participate in the formation of liquid crystals in the presence of a second surfactant. Such a surfactant system forms a layer of lamellar liquid crystals around the primary W_1/O emulsion to provide a barrier between the primary emulsion and the external aqueous W_2 phase. This type of an emulsion is different from conventional emulsions which rely upon the orientation of the hydrophobic and hydrophilic components of a surfactant at an oil-water interface. The formation of a layer of lamellar liquid crystals around the W_1/O primary emulsion can be detected by the presence of Maltese crosses viewed by optical microscopy through crossed polarizing plates or by freeze fracture electron microscopy.

As previously described, the HLB value of a surfactant is an approximate measurement of the solubility of a surfactant in water or oil. However, an HLB value is used only when the concentration of a surfactant in solution is sufficiently low such that a molecular monolayer of the surfactant is formed around droplets dispersed in a continuous phase, i.e., around the primary W_1/O emulsion. At higher surfactant concentrations, the surfactant can undergo a phase transition and the oil droplets then can be surrounded by a layer of lamellar liquid crystals, which can improve emulsion stability. The liquid crystals are localized at the oil/water interface and have been observed by microscopy techniques.

It is theorized, but not relied upon herein, that the layer of lamellar liquid crystals provides improved multiple emulsion stability for the following reasons. As primary W_1/O emulsion droplets approach one another, multiple emulsion instability occurs in two steps: (1) flocculation of the W_1/O droplets due to van der Waals forces, followed by (2) coalescence of the flocculated W_1/O droplets. The rate of droplet coalescence is related to the stability of the emulsifier film surrounding the droplets.

Liquid crystals stabilize emulsions by affecting both steps of the instability process. First, flocculation is reduced because the presence of liquid crystals modifies the attractive van der Waals forces. The presence of liquid crystal layers around the droplets significantly reduces interdroplet attraction. Additionally, the liquid crystals provide mechanical stability to the film around the primary W_1/O droplets, and thereby resist the film-thinning process which leads to droplet coalescence.

Exemplary classes of surfactants capable of participating in the formation of a liquid crystal structure around the primary W_1/O emulsion to stabilize the W_1-O-W_2 multiple emulsion composition include, but are not limited to, quaternary ammonium compounds having two long alkyl chains (i.e., C_8-C_{22}), di-long chain alkyl (C_8-C_{22}) amines, long chain fatty (C_8-C_{22}) alcohols, ethoxylated fatty (C_8-C_{22}) alcohols, double-tailed anionic surfactants, and double-tailed phospholipids. Specific surfactants capable of participating in the formation of a stabilizing liquid crystal structure include, but are not limited to dicyetylmonium chloride, distearylmonium chloride, dipalmitylamine, cetyl alcohol, stearyl alcohol, steareth-2, steareth-21, dioctylsodium sulfosuccinate, phosphatidylserine, phosphatidylcholine and mixtures thereof.

The preferred surfactant systems capable of forming a stabilizing liquid crystal structure are surfactant blends that maintain the layer of lamellar liquid crystals below the Krafft temperature of the surfactants. The Krafft temperature corresponds to the melting point of the hydrocarbon chains in the surfactant, and is the temperature above which the surfactant hydrocarbon chains are liquid-like and disordered. Preferred liquid crystal-forming surfactants include, but are not limited to, a mixture of cetyl alcohol, stearyl alcohol, dicyetylmonium chloride, stearyl amidopropyl dimethylamine and cetareth-20; a mixture of distearylmonium chloride, cetrimonium chloride, cetyl alcohol and stearyl alcohol; and a mixture of oleth-15, cetyl alcohol and stearyl alcohol.

In a preferred embodiment, the external aqueous phase comprises water, an emulsifier and a second topically-active compound. If the W₁-O-W₂ multiple emulsion composition is designed to be a hair shampoo, skin cleanser or rinse-off conditioner, the emulsifier present in the external aqueous phase can serve both as the emulsifier and the second topically-active compound.

The second topically-active compound is present in an amount of 0% to about 30%, and preferably about 0.1% to about 30%, by weight of the external aqueous phase. Alternatively expressed, the second topically-active compound is present in an amount of 0% to about 30%, and preferably about 0.05% to about 30% by weight of the W₁-O-W₂ multiple emulsion composition.

The second topically-active compound typically is a water-soluble compound which performs its intended function, such as cleansing or conditioning, prior to rinsing the W₁-O-W₂ multiple emulsion composition from the skin or hair. The second topically-active compound can be removed from the hair or skin with the remainder of the external aqueous phase during the rinsing step (e.g., a cleansing agent), or can remain on the hair after the rinsing step (e.g., a conditioner that is substantive to the hair).

Therefore, in addition to cleansing agents, the second topically-active compound can be a hair conditioner that is substantive to the hair, such as a quaternary ammonium compound, and therefore is not rinsed from the hair during the rinsing step. The second topically-active compound also can be a reducing agent used in permanent waving, such as ammonium bisulfite, ammonium sulfate, ammonium sulfite, cysteamine, ammonium thioglycolate, cysteine, cysteine hydrochloride, ethanolamine thioglycolate, hydroquinone, glyceryl thioglycolate, mercaptopropionic acid, potassium metabisulfite, potassium sulfite, potassium thioglycolate, sodium bisulfite, sodium hydrosulfite, sodium hydroxymethane sulfonate, sodium metabisulfite, sodium sulfite, sodium thioglycolate, thioglycerin, thioglycolic acid, thiolactic acid, thiosalicylic acid, or mixtures thereof. In addition, a reducing agent can be the first topically-active compound, when the second topically-active compound is a hair cleanser.

The external aqueous phase also can include optional ingredients traditionally included in topically-applied compositions, such as fragrances, antioxidants, detackifying agents, preservatives or dyes. The external aqueous phase may include a nonpolymeric gelling agent or viscosity enhancer, such as an inorganic gelling agent, like fumed silica for example, but is free of polymeric organic gelling agents.

A W₁-O-W₂ multiple emulsion composition of the present invention therefore is free of polymeric gelling agents in the external aqueous phase, and can be a liquid composition having a viscosity of about 1 to about 15,000 cps, as measured on a Brookfield Viscometer using a #4 spindle at a speed of 20 rpm. For a hair care composition, the composition preferably has a viscosity of about 100 to about 10,000 cps. To achieve the full advantage of the present invention, the W₁-O-W₂ multiple emulsion composition has a viscosity of about 1,000 to about 9,000 cps.

The W₁-O-W₂ multiple emulsion also can be a cream-like composition having a viscosity of about 50,000 to about 1,200,000 cps, as measured on a Brookfield Viscometer using a T-BAR spindle at a speed of 2.5 rpm. The relatively high viscosity can be achieved by the addition of a nonpolymeric gelling agent or can arise due to the identity and amount of ingredients in the W₁-O-W₂ multiple emulsion composition.

The W₁-O-W₂ multiple emulsion compositions typically are prepared by a two-step process. In the two-step process, the primary water-in-oil (W₁/O) emulsion is manufactured first, then is dispersed in the external aqueous phase (W₂). The second step of the two-step process emulsifies the primary W₁/O emulsion. This two-step manufacturing method allows for the preparation of an internal aqueous phase (W₁) different in composition from the external aqueous phase (W₂).

In the manufacture of the primary W₁/O emulsion, the surfactant phase is admixed with the oil phase, then the aqueous phase is dispersed in the combined surfactant and oil phases under high shear conditions and preferably in a homogenizer (e.g., about 17,000 rpm in an IKA Homogenizer, available from IKA Laboratories, Cincinnati, OH). Primary emulsions that were homogenized provided smaller particle size droplets of the internal aqueous phase (W₁) (i.e., about 1 μ to about 75 μ in diameter), which were more uniform in size. Primary emulsions that were homogenized therefore had increased stability.

The resulting primary W₁/O emulsion then is dispersed in the external aqueous phase (W₂) using low shear mixing (e.g., about 130 rpm on a bench scale mixer). Intensive, high shear mixing during this manufacturing step has a tendency to destroy the primary emulsion and release the contents of the internal W₁ aqueous phase.

The two-step manufacturing method also allows a formulator to prepare different primary W₁/O emulsions, i.e., primary emulsions that incorporate different topically-active compounds into the internal aqueous phase. Then, the W₁/O primary emulsions each can be converted into a W₁-O-W₂ multiple emulsion composition having an external aqueous phase (W₂) of the same composition. The two W₁-O-W₂ compositions then can be admixed to provide a multiple emulsion composition incorporating different primary W₁/O emulsions having different topically-active compounds which perform different or complementary functions.

The following examples are illustrative of the primary W₁/O emulsions and the W₁-O-W₂ multiple emulsion compositions of the present invention. However, the present invention is not limited to the specific examples set forth below. In the following examples, all amounts of the various ingredients are expressed by weight percentages unless otherwise specified.

As demonstrated in the following examples, the W₁-O-W₂ multiple emulsion compositions of the present invention were phase-stable for at least a sufficient time to detect the W₁-O-W₂ multiple emulsion by an optical microscope. It is within the skill of persons in the art to optimize the identity of composition ingredients and weight percentages to provide W₁-O-W₂ multiple emulsion compositions that are phase-stable over the expected life of the composition. The W₁-O-W₂ multiple emulsion compositions also were moderately viscous liquids to creams; were easy to apply; and effectively delivered the topically-active compounds to the skin or hair. Each of the following examples was prepared by the above-described two-step method.

The following Examples 1-30 illustrate W₁-O-W₂ multiple emulsion compositions wherein the first topically-active compound in the internal W₁ aqueous phase is a hair fixative resin and the second topically-active compound in the external W₂ aqueous phase is a cationic hair conditioner or an anionic hair cleanser. The cationic surfactant systems present in the external aqueous phase (W₂) of Examples 1-28 and 30 form a layer of lamellar liquid crystals around the primary W₁/O emulsion to stabilize the W₁-O-W₂ multiple emulsion compositions. The anionic surfactants present in the composition of Example 29 stabilize the emulsion.

In addition to Examples 1-30, numerous other combinations of first and second topically-active compounds are envisioned. For example, if the second topically-active compound is a conditioner, the first topically-active compound can be a true bodifier (e.g., a styling aid resin), a curling iron-activated styling aid, a protein (e.g., hydrogenated wheat protein or hydrolyzed keratin protein), a sunscreen, an optical brightener (e.g., 2-pyrrolidone-5-carboxylic acid), a humectant, or a water-soluble or water-dispersible hair dye.

If the second topically-active ingredient is a skin cleanser, the first topically-active ingredient can be a skin care product for dry skin (e.g., glycerin, propylene glycol, sorbic acid, proline, urea, hyaluronate of dimethylsilanol, allantoin, aloe vera, pyrrolidone carboxylic acid or hydrolyzing collagen) or for oily skin (e.g., provitamin B₅).

In accordance with an important feature of the present invention, incompatible compounds can be incorporated as the first and second topically-active compounds because the incompatible compounds do not come in contact in the W₁-O-W₂ multiple emulsion composition. For example, if the second topically-active compound is a cleanser (e.g., an anionic surfactant), the first topically-active compound can be a hair conditioner (e.g., a cationic surfactant), a hair fixative, a sunscreen, an optical brightener, a hair dye, a deodorant, a skin care product, a humectant, or a bactericide or other topical medicament.

Exemplary, but non-limiting, combinations of first and second topically-active compounds in a W₁-O-W₂ multiple emulsion composition of the present invention are:

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	First Topically-Active Compound	Second Topically-Active Compound
5	Neutralizer for reducing agents	Reducing agent for permanent wave
	Reducing Agent for Permanent Wave	Hair Cleanser
	Reducing Agent for Permanent Wave	Hair Conditioner
10	Neutralizer	Cysteamine
	Neutralizer	Hair Conditioner
	Hair Conditioner	Neutralizer
15	Hair Fixative Resin	Hair Fixative Resin (incompatible)
	Topical Medicament	Topical Anesthetic
	Skin Conditioner	Make-up Remover
	Sunburn Reliever (e.g.,vitamin F)	Skin Cleanser
20	Sunscreen	Skin Cleanser
	Skin Conditioner	Skin Cleanser
	Bacteriocide	Skin Cleanser
25	Deodorant	Skin Cleanser
	Antifungal Compound	Skin Cleanser
	Anesthetic	Skin Cleanser
	Anti-Inflammatory	Skin Cleanser
30	Anti-Irritant	Skin Cleanser
	Medicament	Skin Cleanser
	Anti-Itch Compound	Skin Cleanser
35	Self-Tanning Compound	Skin Cleanser
	Self-Tanning Compound	Skin Conditioner
	Dermatitis Medication	Skin Cleanser
	Hair Conditioner	Hair Cleanser
40	Hair Fixative	Hair Cleanser
	Hair Dye	Hair Cleanser
	Hair Dye	Hair Conditioner
45	Hair Dye	Bleaching Agent
	Sunscreen	Hair Conditioner
	Protein	Hair Conditioner
	Humectant	Hair Conditioner
50	Hair Growth Promoter	Hair Cleanser
	Hair Bleaching Agent	Hair Cleanser
	Water-Soluble Hair Conditioner	Substantive Hair Conditioner

EXAMPLES 1-30

Examples 1-30 were prepared in an essentially identical manner by the two-step process. The primary W₁/O emulsion first was prepared as described above. The primary emulsion of each example had a ratio of aqueous phase/oil phase/surfactant phase (W/O/S) summarized in Table 1. The aqueous phase W₁/O incorporated about 20% to about 40% by weight of the primary emulsion of a hair fixative resin, such as polyvinylpyrrolidone or the quaternized copolymer of vinylpyrrolidone and dimethylaminoethyl methacrylate, as summarized in Table 1. The surfactant phase was either cetyl dimethicone copolyol or dimethicone copolyol. The oil phase was a volatile silicone compound, i.e., cyclomethicone. It was found that primary emulsions containing about 60% by weight aqueous phase W₁ had excellent stability.

In Examples 1-28, the external aqueous phase (W₂) was Hair Conditioner Base No. 1 having the following formula:

Hair Conditioner Base No. 1 (External Aqueous Phase)	
Ingredient	Weight % ⁵⁾
Soft water	q.s. to 100
Citric acid (50% aqueous)	0.185
Stearamidopropyltrimethylamine ¹⁾	0.5
Propylene glycol	0.5
Dicetyldimonium chloride/PG ²⁾	2.1
Stearyl alcohol and Ceteareth-20 Blend ³⁾	1.0
Cetyl alcohol	3.25
Potassium chloride	0.1
Disodium EDTA	0.1
Methylchloroisothiazoline/methylisothiazolinone ⁴⁾	0.05
DMDM hydantoin ⁴⁾	0.1
Potassium hydroxide (50% aqueous)	0.033
Fragrance	0.2

¹⁾ LEXAMINE S-13, available as a 100% active material, from Inolex Chemical Co., Philadelphia, PA;

²⁾ VARISOFT 432PG, available as a 68% active material, from Sherex Chemical Co., Dublin, OH, to provide 1.43% active material in the final formulation;

³⁾ PROMULGEN G, available as a 100% active blend, from Amerchol Corp., Edison, NJ;

⁴⁾ preservatives; and

⁵⁾ based on the weight raw material added to the hair conditioner base.

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In Example 29, the external aqueous phase was Hair Cleanser Base No. 1 having the following formula:

5 Hair Cleanser Base No. 1 ⁹⁾ (External Aqueous Phase)	
Ingredient	Weight % ⁶⁾
Ammonium Lauryl Sulfate ⁷⁾	3.75
Sodium Lauryl Sulfate ⁸⁾	12.75
Ammonium Chloride	2.5
Soft Water	q.s. to 100

⁶⁾ based on the weight of active ingredients;

⁷⁾ added as a 30% active aqueous solution;

⁸⁾ added as a 30% active aqueous solution; and

⁹⁾ includes 16.5% total anionic surfactant as a hair cleanser.

In Example 30, the external aqueous phase was Hair Conditioner Base No. 2 having the following formula:

25 Hair Conditioner Base No. 2 (External Aqueous Phase)	
Ingredient	Wt. % ⁵⁾
Soft water	q.s. to 100
Citric acid (50% aqueous)	0.02
Hydroxyethylcellulose	0.3
Distearyldimonium chloride ¹⁰⁾	1.1
Cetrimonium chloride ¹¹⁾	1.2
Cetearyl alcohol ¹²⁾	1.9
Dyes	0.0008
Potassium chloride	0.3
Disodium EDTA	0.01
Methylchloroisothiazoline/methylisothiazolinone ⁴⁾	0.05
DMDM hydantoin ⁴⁾	0.2
Fragrance	0.2

¹⁰⁾ AROSUFT® TA-100, available as a 100% active material, from Witco Corp., NY, NY;

¹¹⁾ BARQUAT® CT-29, available as a 30% active material, from Lonza, Inc., Fairlawn, NJ, to provide 0.57% active material in the final formulation; and

¹²⁾ available as a 100% active material, from Witco Corp., NY, NY.

The primary W₁/O emulsion was added to the external phase, with moderate stirring, in an amount of 4.5% to 55% by weight, based on the final weight of the W₁-O-W₂ multiple emulsion composition.

In particular, the W₁-O-W₂ multiple emulsion compositions of Examples 1-28 were manufactured as follows. Hair Conditioner Base No. 1 was prepared by heating the soft water (i.e., tap water softened by passage through an ion exchange column) to about 155 to 160°F, then adding the citric acid, stearamidopropyltrimethylamine, and propylene glycol to the heated water. Next, the dicetylmonium chloride/PG was added, and the resulting mixture was stirred for 5 minutes. When the mixture was homogeneous, the stearyl alcohol/ceteareth-20 blend was added to the mixture at a

temperature above 145°F. The cetyl alcohol was added next, and the resulting mixture was stirred at high speed (i.e., about 130 rpm on a production scale mixer). After all the ingredients were dispersed, mixing was continued for 30 minutes at 155°F. Then, the mixture was cooled to 130-135°F, followed by the addition of potassium hydroxide (diluted 1:8 with soft water). In a separate container, the potassium chloride and disodium EDTA were completely dissolved in 120°F soft

5 water, then, at 117-122°F, the salt solution was slowly added to the batch mixture. After cooling the resulting mixture, i.e., Hair Conditioner Base No. 1, to 110°F or less, the primary W₁/O emulsion, preservatives and fragrance were added to the mixture. The resulting W₁-O-W₂ multiple emulsion composition was cooled to 100-105°F, and low shear mixing was continued for about 30 minutes. Finally, the pH was adjusted to about 4.5 to about 6.1 with liquid citric acid or potassium hydroxide. Alternatively, the primary W₁/O emulsion can be incorporated into the mixture after pH adjustment.

10 The compositions of Examples 29 and 30 were prepared in a similar two-step method.

The following Table I illustrates Examples 1-30 wherein different parameters with respect to the primary W₁/O emulsion and the external aqueous phase (W₂) were varied. Examples 1 and 2 are control examples in which cyclomethicone was incorporated directly into the hair conditioner base. The presence of a W₁-O-W₂ multiple emulsion was confirmed by optical microscopy.

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TABLE I

EXAMPLE	% SILICONE ¹⁹	W/O/S ¹⁴	SURFACTANT ¹⁹	RESIN ¹⁹	REMARKS
1 (comparative)	1.8	--	--	--	Hair Conditioner Base No. 1 (control)
2 (comparative)	3.0	--	--	--	Hair Conditioner Base No. 1 with more cyclomethicone (control)
3	1.8	46/46/8	Cetyl Dimethicone Copolyol ¹⁴	--	Multiple emulsion observed under a light microscope
4	1.8	60/30/10H ²⁰	Cetyl Dimethicone Copolyol ¹⁴	Polyvinylpyrrolidone ¹⁹ (1.1 %)	Curl retention (qualitative) was noted on hair tresses, resin coating felt on hair
5	3.0	60/30/10H	Cetyl Dimethicone Copolyol ¹⁴	Polyvinylpyrrolidone ¹⁹ (1.8 %)	Resin coating felt on hair; stable for 3 months ²¹ ; weak set retention of tresses ²²
6	4.0	60/30/10H	Cetyl Dimethicone Copolyol ¹⁴	Polyvinylpyrrolidone ¹⁹ (2.4 %)	Identical set retention (qualitative) as Example 5
7 (comparative)	3.0	--	Cetyl Dimethicone Copolyol ¹⁴	Polyvinylpyrrolidone ¹⁹	Components added separately, no primary emulsion formed; set retention less than Example 5
8	4.5	60/30/10H	Cetyl Dimethicone Copolyol ¹⁴	Polyvinylpyrrolidone ¹⁹ (2.7 %)	Tress curl retention improved over Example 5
9	3.0	50/40/10	Cetyl Dimethicone Copolyol ¹⁴	Quaternized copolymer of vinylpyrrolidone and dimethylaminoethyl methacrylate (Polyquaternium 11) ²⁰ (0.75 %)	Set retention greater than polyvinylpyrrolidone (qualitative) of Example 5; stable for 3 months
10	4.0	50/40/10	Cetyl Dimethicone Copolyol ¹⁴	Quaternized copolymer of vinylpyrrolidone and dimethylaminoethyl methacrylate (Polyquaternium 11) ²⁰ (0.90 %)	Slightly better set retention than Example 9 (qualitative); stable for 3 months
11	3.0	70/20/10UH	Cetyl Dimethicone Copolyol ¹⁴	Quaternized copolymer of vinylpyrrolidone and dimethylaminoethyl methacrylate (Polyquaternium 11) ²⁰ (2.1 %)	Salon test ²³ ; wetcomb poorer than control (Example 1) hold and set retention identical to control performance: curl retention significantly better than control; stable for 3 months

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TABLE I					
EXAMPLE	% SILICONE ¹¹	W/O/S ¹⁴	SURFACTANT ¹³	RESIN ¹⁵	REMARKS
12	3.0	70/20/10H	Cetyl Dimethicone Copolyol ¹⁶	Quaternized copolymer of vinylpyrrolidone and dimethylaminooethyl methacrylate (Polyquaternium 11) ²⁰ (2.1 %)	Salon test: weicomb poorer than control hold and set retention greater than control performance: curl retention significantly better than Example 11; stable for 3 months
13 (comparative)	3.0	70/20/10	Cetyl Dimethicone Copolyol ¹⁶	Quaternized copolymer of vinylpyrrolidone and dimethylaminooethyl methacrylate (Polyquaternium 11) ²⁰ (2.1 %)	Components added separately sensory performance poorer than Example 14
14	3.0	70/20/10H	Cetyl Dimethicone Copolyol ¹⁶	Quaternized copolymer of vinylpyrrolidone and dimethylaminooethyl methacrylate (Polyquaternium 11) ²⁰ (2.1 %)	Sensory performance greater than Example 13
15 (comparative)	3.0	80/17/3	Dimethicone copolyol ¹⁷	Quaternized copolymer of vinylpyrrolidone and dimethylaminooethyl methacrylate (Polyquaternium 11) ²⁰ (2.1 %)	Components added separately salon: observed poorer performance compared to Ex. 16
16	3.0	80/17/3H	Dimethicone copolyol ¹⁷	Quaternized copolymer of vinylpyrrolidone and dimethylaminooethyl methacrylate (Polyquaternium 11) ²⁰ (2.8 %)	Salon test: no difference in hold/curl retention compared to Ex. 9, stable for about 2 months
17 (comparative)	1.8	80/17/3	Dimethicone copolyol ¹⁷	Quaternized copolymer of vinylpyrrolidone and dimethylaminooethyl methacrylate (Polyquaternium 11) ²⁰ (1.7 %)	Components added separately
18	1.8	80/17/3H	Dimethicone copolyol ¹⁷	Quaternized copolymer of vinylpyrrolidone and dimethylaminooethyl methacrylate (Polyquaternium 11) ²⁰ (1.7 %)	Added as emulsion, significant increase in composition viscosity compared to Example 17
19 (comparative)	1.8	85/10/5	Dimethicone copolyol ¹⁷	Quaternized copolymer of vinylpyrrolidone and dimethylaminooethyl methacrylate (Polyquaternium 11) ²⁰ (3.1 %)	Components added separately

TABLE I

EXAMPLE	% SILICONE ^a	W/O/S ^a	SURFACTANT ^b	RESIN ^c	REMARKS
20	1.8	85/10/SH	Dimethicone copolyol ^d	Quaternized copolymer of vinylpyrrolidone and dimethylaminoethyl methacrylate (Polyquaternium 11) ^e (3.1 %)	Viscosity 6000 cps, salon study 4 of 6 models had worse curl retention than identical composition using cetyl dimethicone copolyol (Example 21), stability about 45 days
21	1.8	85/10/SH	Cetyl Dimethicone Copolyol ^d	Quaternized copolymer of vinylpyrrolidone and dimethylaminoethyl methacrylate (Polyquaternium 11) ^e (3.1 %)	Viscosity 1080 cps, stability greater than 45 days
22	3.0	85/10/5 H	Dimethicone Copolyol ^d	Quaternized copolymer of vinylpyrrolidone and dimethylaminoethyl methacrylate (Polyquaternium 11) ^e (5.1 %)	Salon study: worse combing but better curl than control Example 1; stability greater than one month
23	3.0	85/10/5 H	Cetyl Dimethicone Copolyol ^d	Quaternized copolymer of vinylpyrrolidone and dimethylaminoethyl methacrylate (Polyquaternium 11) ^e (5.1 %)	Salon study: worse combing, but better curl retention after 4 to 5 hours; stability greater than one month
24	5.0	85/10/5 H	Cetyl Dimethicone Copolyol ^d	Quaternized copolymer of vinylpyrrolidone and dimethylaminoethyl methacrylate (Polyquaternium 11) ^e (8.5 %)	Stability greater than 12 days
25	5.5	85/10/5 H	Cetyl Dimethicone Copolyol ^d	Quaternized copolymer of vinylpyrrolidone and dimethylaminoethyl methacrylate (Polyquaternium 11) ^e (9.35 %)	Stability greater than 2 days
26	1.8	87/10/3 H	Dimethicone Copolyol ^d	Quaternized copolymer of vinylpyrrolidone and dimethylaminoethyl methacrylate (Polyquaternium 11) ^e (3.1 %)	Stability about 45 days
27	3.0	70/20/10 H	Cetyl Dimethicone Copolyol ^d	Vinylpyrrolidone/methacrylamido-propyltrimethylammonium chloride copolymer (Polyquaternium 28) ^e (2.1 %)	Stability greater than 7 days

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TABLE I					
EXAMPLE	% SILICONE ¹⁹	W/O/S ²⁰	SURFACTANT ²¹	RESIN ¹⁹	REMARKS
28	3.0	70/20/10 H	Cetyl Dimethicone Copolyol ¹⁹	PVP/Dimethylaminoethyl-methacrylate Copolymer ²⁰ (2.1 %)	Stability greater than 7 days
29 ²²	2.0	85/10/5 H	Cetyl Dimethicone Copolyol ¹⁹	Quaternized copolymer of vinylpyrrolidone and dimethylaminoethyl methacrylate (Polyquaternium 11) ²⁰ (3.4 %)	A 3-in-1 shampoo/conditioner/styling aid; stability about 5 minutes
30 ²³	3.0	85/10/5 H	Cetyl Dimethicone Copolyol ¹⁹	Quaternized copolymer of vinylpyrrolidone and dimethylaminoethyl methacrylate (Polyquaternium 11) ²⁰ (5.1 %)	Treated tresses exhibited extra body in salon tests; stability greater than 5 days

- (19) % volatile silicone (cyclomethicone) in the final W₁-O-W₂ multiple emulsion composition or control;
- (20) weight ratio of water phase (W) to volatile oil phase (O) to surfactant phase (S) in the primary W₁/O emulsion;
- (21) surfactant present in the surfactant phase of the primary W₁/O emulsion;
- (22) ABIL EM-90, available as a 100% active compound from Goldschmidt Chemical Corporation, Hopewell, VA;
- (23) SILWET Surfactant L-7622, available as a 100% active compound from OSI Specialties, Inc., Tarrytown, NY;
- (24) Hair fixative resin present in the primary W₁/O emulsion as the first topically-active compound, and the % by weight of the hair fixative resin in the W₁-O-W₂ multiple emulsion composition;
- (25) PVP K-30 (M.W. 38,000), available from ISP Corp., Wayne, NJ, and added as a 30% aqueous solution;
- (26) GAFQUAT 755N (M.W. 1,000,000), available from ISP Corp., Wayne, NJ, as a 20% aqueous solution;
- (27) H: homogenized;
UH: unhomogenized;
- (28) sample stability tested at room temperature;
- (29) comparison to control sample of Example 1 unless otherwise noted;
- (30) salon tests were performed on six models;

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- 50 GAFQUAT HS-100 (M.W. 1,000,000), available from ISP Chemicals, Inc., Wayne, NJ, as a 20% aqueous solution;
- 55 COPOLYMER 845 (M.W. 1,000,000), available from ISP Chemicals, Inc., Wayne, NJ, as a 20% aqueous solution;
- 60 *m* the external aqueous phase was the anionic surfactant-based Hair Cleanser Base No. 1; and
- 65 *n* the external aqueous phase was the cationic surfactant-based Hair Conditioner Base No. 2.

Various compositions summarized in Table I were applied to virgin brown hair tresses, available from DeMeo Bros., NY, NY. The compositions were worked into the hair, then rinsed from the hair. The hair tresses then were dried. The hair tresses were examined visually. Salon tests on six human models also were conducted. The technicians empirically rated hair treated with the W₁-O-W₂ multiple emulsion compositions, and the results were averaged.

As shown in Table I, the W₁-O-W₂ multiple emulsion compositions outperformed the comparative compositions of Examples 1, 2, 7, 13, 15, 17 and 19, wherein the ingredients were added separately as opposed to first forming a primary W₁/O emulsion.

In particular, mass spectrometry indicated that cyclomethicone remained on the hair after rinsing, i.e., the primary emulsion was deposited. After drying the hair, the resin was released to impart set retention properties to the hair.

In addition, it was observed that resin was deposited on the hair. Hair treated with the compositions of Examples 11 and 12 had detectable amounts of resin deposited on the hair shaft. This was confirmed by scanning electron micrographs of hair treated with the compositions of Examples 11 and 12, compared to hair treated with a control sample (Example 2) that did not include a resin and compared to rinsed and unrinsed hair samples treated with an aqueous

resin solution (0.5% by weight GAFQUAT 755N). The composition of Example 12, which was homogenized, applied a more uniform and thinner coating of resin on the hair, which is preferred.

Table I also illustrates that the W₁-O-W₂ multiple emulsion compositions of the present invention exhibit a stability of greater than 3 months (Examples 5 and 9-12) and that 55% by weight of a primary W₁/O emulsion can be incorporated into the W₁-O-W₂ multiple emulsion composition (Example 25).

In addition to the compositions of Examples 1-30, other W₁-O-W₂ multiple emulsion compositions were prepared which incorporated a first topically-active ingredient different from a hair styling resin. These W₁-O-W₂ compositions are illustrated in Table II as Examples 31-41. The compositions of Examples 31-41 incorporated either a water-soluble dye, a sunscreen, an optical brightener or a permanent wave composition as the first topically-active compound in the primary W₁/O emulsion. The compositions of Examples 31-41 were prepared by the two-step method set forth above with respect to Examples 1-30.

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TABLE II

EXAMPLE	% SILICONE ¹¹	W/O/S ¹⁴	SURFACTANT ¹⁵	FIRST TOPICALLY ACTIVE COMPOUND	EXTERNAL PHASE	REMARKS
31	1.87	72/18.7/9.3	Cetyl Dimethicone Copolyol ¹⁶	FD&C Red No. 22 (dye) 10% ¹⁷	Hair Conditioner Base No. 1	The composition was applied to virgin white tresses of hair. The composition of Example 31 dyed the tresses. No dye deposition observed with a control oil-in-water composition; stability at least one day ¹⁸
32	1.87	72/18.7/9.3	Cetyl Dimethicone Copolyol ¹⁶	Tri-K Arianor Madden Red (dye) 1% ¹⁹	Hair Conditioner Base No. 1	Stability greater than 6 days at room temperature and at 110° F, up to 6 days at 120° F
33	1.87	72/18.7/9.3	Cetyl Dimethicone Copolyol ¹⁶	Tri-K Arianor Madden Red (dye) 1% and Tri-K Arianor Steel Blue (dye) 1% ¹⁹	Hair Conditioner Base No. 1	Individual W ₁ droplets including either blue or red dye were observed under an optical microscope
34	3.44	74/20.7/5.3	Cetyl Dimethicone Copolyol ¹⁶	Benzophenone-4 (sunscreen) 5% ²⁰	Hair Conditioner Base No. 1	The presence of sunscreen on treated hair was detected by UV/visible light absorption; gas chromatography/mass spectroscopy detected about 10 ppm of sunscreen per gram of hair; stability greater than one week
35	2.76	77.7/13.8/8.5	Cetyl Dimethicone Copolyol ¹⁶	Stilbene Derivative ²¹ (optical brightener) 10% ²²	Hair Conditioner Base No. 1	Stability at least one week
36	13.5	70/20/10	Cetyl Dimethicone Copolyol ¹⁶	hydrogen peroxide (permanent wave neutralizer) ²³ 1.54% ²⁴	Hair Reducing Base No. 1 ¹⁹ (90%) and Dimethicone Copolyol ¹⁶ (10%)	A W ₁ -O-W ₂ multiple emulsion was observed, but less than 100% formation; stability about 4 hours
37	13.5	70/20/10	Cetyl Dimethicone Copolyol ¹⁶	hydrogen peroxide (permanent wave neutralizer) ²³ 1.54% ²⁴	Hair Reducing Base No. 1 ¹⁹ (90%) and Dimethicone Copolyol ¹⁶ (10%)	A W ₁ -O-W ₂ multiple emulsion was observed, but less than 100% formation; stability about 4 hours

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TABLE II

EXAMPLE	% SILICONE ^a	W/O/S ^b	SURFACTANT ^c	FIRST TOPICALLY-Y. ACTIVE COMPOUND	EXTERNAL PHASE	REMARKS
38	20.0	63/27/10	Cetyl Dimethicone Copolyol ^d	hydrogen peroxide (permanent wave neutralizer) ^e 1.39% ^f	Hair Reducing Base No. 1 ^g (91%) and Dimethicone Copolyol ^h (9%)	A W ₁ -O-W ₂ multiple emulsion was observed, but less than 100% formation; stability about 2 hours
39	31.5	45/45/10	Cetyl Dimethicone Copolyol ⁱ	hydrogen peroxide (permanent wave neutralizer) ^j 1.00% ^k	Hair Reducing Base No. 1 ^l (90%) and Dimethicone Copolyol ^m (10%)	A W ₁ -O-W ₂ multiple emulsion was observed, but less than 100% formation; stability about 2 hours
40	21.0	60/30/10	Cetyl Dimethicone Copolyol ⁱ	hydrogen peroxide (permanent wave neutralizer) ^j 1.32% ^k	Hair Reducing Base No. 2 ^l (90%) and Dimethicone Copolyol ^m	A W ₁ -O-W ₂ multiple emulsion was observed, but less than 100% formation; stability about 2 hours
41	3.0	70/20/10	Cetyl Dimethicone Copolyol ⁱ	Hair Reducing Base No. 2 (5.75% cysteamine HCl 0.5% ammonium hydroxide 1.4% ammonium chloride)	Hair Conditioner Base No. 1	Stability greater than 3 weeks

^a ABIL[®] Wax 2434, available as a 100% active compound from Goldschmidt Chemical Corp., Hopewell, VA.;^b wt. % of first topically-active material in W₁;^c LEUCOPHOR BSB, available as a 100% active compound from Sandoz Chemicals Corp., Charlotte, NC;^d permanent wave neutralizer including 2.2% (by weight) hydrogen peroxide, 0.005% phosphoric acid and the remainder water;

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The compositions of Examples 31 and 34 demonstrate W_1 -O- W_2 multiple emulsion compositions of the present invention that incorporate incompatible first and second topically-active compounds. In Example 31, the first topically-active compound is an anionic dye and the second topically-active compound is a cationic quaternary ammonium compound. Simply admixing aqueous solutions of the anionic dye and cationic quaternary ammonium compound resulted in an immediate precipitate. However, adding the primary W_1 /O emulsion incorporating the anionic dye to the external aqueous phase (W_2) incorporating the cationic salt provided a homogeneous composition, without the formation of a precipitate. The primary W_1 /O emulsion prevents formation of a precipitate because the oil phase separates the incompatible topically-active ingredients.

The composition of Example 31 also effectively dyed virgin white tresses of hair. White hair tresses were not dyed by a composition wherein the anionic dye was simply admixed with the cationic quaternary ammonium compound. In this composition, the dye was not available to treat the hair because the dye interacted with the cationic quaternary ammonium compound.

Similarly, Example 34 incorporates an anionic sunscreen agent and a cationic quaternary ammonium salt. If benzophenone-4 is added directly to Hair Conditioner Base No. 1, a precipitate results. However, adding a primary W_1 /O emulsion incorporating benzophenone-4 into the internal W_1 phase prevented formation of a precipitate. In addition, the compositions of Examples 36-40 illustrate that an oxidizing agent (e.g., hydrogen peroxide) can be combined with an incompatible reducing agent in a W_1 -O- W_2 multiple emulsion composition.

The following Table III illustrates other primary W_1 /O emulsions that have been prepared, then incorporated into Hair Conditioner Base No. 1 to provide a W_1 -O- W_2 multiple emulsion composition of the present invention. Hair Condi-

	31	Hair Reducing Base No. 1 includes:	
		water	67.4 (wt. %)
		ammonium thioglycolate (60% solution)	23.0
		dithioglycolate (40% solution)	5.0
		ammonium hydroxide (28%)	2.1
		ammonium bicarbonate	2.0
		glycenn	0.5;
	32	AMERSIL DMC357, available as a 100% active compound from Amerchol Corp., Edison, NJ;	
	33	DC 193, available as a 100% active compound from Dow Corning Corp., Midland, MI;	
	34	Hair Reducing Base No. 2 includes:	
		water	84.5 (wt. %)
		cysteamine HCl (75%)	11.0
		ammonium hydroxide (28%)	2.5
		ammonium bicarbonate	2.0; and
	35	ABIL [®] B8863, available as a 100% material from Goldschmidt Chemical Corp., Hopewell, VA.	

tioner Base No. 1 is a cationic composition capable of emulsifying silicone oils and destroying the primary emulsion. However, the primary W/O emulsions were stable when emulsified into the cationic hair conditioner base under low shear conditions. Examples 1-95 also provided evidence that the water-soluble, first topically-active compound present in the internal aqueous phase remains on the hair in sufficient quantity to perform its intended function.

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TABLE III

EXAMPLE	W/O/S ¹⁴	% SILICONE ²⁵	SURFACTANT ¹⁵	INTERNAL AQUEOUS PHASE ²⁶
42	70/20/10	20	Cetyl Dimethicone Copolyol ¹⁶	PVP/Dimethylaminoethyl-methacrylate Copolymer ²⁷
43	70/20/10	20	Cetyl Dimethicone Copolyol ¹⁶	Sodium Polystyrene Sulfonate ²⁸
44	85/10/5	10	Dimethicone Copolyol ¹⁷ (60%) and Laureth-4 (40%)	Polyquaternium 11 ²⁹
45	85/10/5	10	Dimethicone Copolyol (60%) and Laureth-1 (40%)	Polyquaternium 11 ²⁹
46	70/20/10	20	Cetyl Dimethicone Copolyol ¹⁶	Polyquaternium 28 ²⁹
47	60/30/10	30	Cetyl Dimethicone Copolyol ¹⁶	Polyvinylpyrrolidone ¹⁹
48	70/20/10	20	Cetyl Dimethicone Copolyol ¹⁶	Polyvinylmethacrylate methyl acrylate decadiene crosspolymer ⁴¹
49	78/16/6	16	Dimethicone Copolyol ¹⁷	Cysteamine soln., 37.5% active
50	73/22/5	22	Cetyl Dimethicone Copolyol ¹⁶	Cysteamine soln., 37.5% active
51	70/20/10	20	Cetyl Dimethicone Copolyol ¹⁶	Cysteamine, 8.25% active
52	45/45/10	45	Cetyl Dimethicone Copolyol ¹⁶	Hair Reducing Base No. 1
53	45/45/10	45	Dimethicone Copolyol ¹⁷	Hair Reducing Base No. 1
54	63/27/10	27	Dimethicone Copolyol ¹⁷	Hair Reducing Base No. 1
55	63/27/10	27	Cetyl Dimethicone Copolyol ¹⁶	Hair Reducing Base No. 1
56	45/45/10	45	Cetyl Dimethicone Copolyol ¹⁶	Hair Reducing Base No. 1
57	45/45/10	45	Dimethicone Copolyol ¹⁷	Hair Oxidizing Base (2.2% hydrogen peroxide)
58	63/27/10	27	Dimethicone Copolyol ¹⁷	Hair Oxidizing Base (2.2% hydrogen peroxide)
59	63/27/10	27	Cetyl Dimethicone Copolyol ¹⁶	Hair Oxidizing Base (2.2% hydrogen peroxide)
60	70/20/10	20	Cetyl Dimethicone Copolyol ¹⁶	Hair Oxidizing Base (2.2% hydrogen peroxide)
61	70/20/10	20	Cetyl Dimethicone Copolyol ¹⁶	1% aqueous red dye solution (Tri-K Arianor Madder Red)
62	70/20/10	20	Cetyl Dimethicone Copolyol ¹⁶	1% aqueous yellow dye solution (Tri-K Arianor Straw Yellow)

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TABLE III				
EXAMPLE	W/O/S ¹⁴	% SILICONE ¹⁵	SURFACTANT ¹⁶	INTERNAL AQUEOUS PHASE ¹⁷
63	70/20/10	20	Cetyl Dimethicone Copolyol ¹⁸	1% aqueous blue dye solution (Tri-K Arianor Steel Blue)
64	70/20/10	20	Dimethicone Copolyol ¹⁷	1% aqueous red dye solution (Tri-K Arianor Madder Red)
65	70/20/10	20	Dimethicone Copolyol ¹⁷	1% aqueous yellow dye solution (Tri-K Arianor Straw Yellow)
66	82/10/8	10	Cetyl Dimethicone Copolyol ¹⁸ (50%) and Stearoxy Dimethicone (50%) ¹⁹	FD&C Blue #1 Dye (5%)
67	72/18.7/9.3	18.7	Cetyl Dimethicone Copolyol ¹⁸	FD&C Blue #1 Dye (10%)
68	72/18.7/9.3	18.7	Cetyl Dimethicone Copolyol ¹⁸	FD&C Red #22 Dye (10%)
69	7/2/1	20	Cetyl Dimethicone Copolyol ¹⁸	1% aqueous sunscreen solution (Benzophenone-4)
70	74/17.2/8.8	17.2	Cetyl Dimethicone Copolyol ¹⁸ (60%) and Stearoxy Dimethicone (40%) ¹⁹	5% aqueous sunscreen solution (benzophenone-4)
71	82.5/10.8/6.7	10.8	Cetyl Dimethicone Copolyol ¹⁸	5% aqueous sunscreen solution (benzophenone-4)
72	77.7/13.8/8.5	13.8	Cetyl Dimethicone Copolyol ¹⁸	10% aqueous optical brightener solution (Stilbene Derivative) ²⁰
73	78.5/14/7.5	14	Dimethicone Copolyol ¹⁷	5% aqueous protein solution (hydrolyzed keratin) ²¹
74	80.2/13.5/6.3	13.5	Cetyl Dimethicone Copolyol ¹⁸	25% aqueous humectant solution (sodium PCA) ²²
75	81.5/10.4/8.1	10.4	Cetyl Dimethicone Copolyol ¹⁸ (55%) and Dimethicone Copolyol ¹⁸ (45%)	25% aqueous humectant solution (sodium PCA) ²²
76	87/9.8/3.2	9.8	Cetyl Dimethicone Copolyol ¹⁸	25% aqueous humectant solution (sodium PCA) ²²
77	80.5/13.3/6.2	13.3	Cetyl Dimethicone Copolyol ¹⁸	50% humectant solution (glycerin)
78	78/14.5/7.5	14.5	Cetyl Dimethicone Copolyol ¹⁸	10% humectant solution (glycolic acid)
79	78.7/14.2/7.1	14.2	Cetyl Dimethicone Copolyol ¹⁸	aqueous phase include 2% FD&C Red #40 dye and 13% Polyquaternium-11 hair fixative resin ²³

TABLE III

EXAMPLE	W/O/S ^{1a}	% SILICONE ^{2b}	SURFACTANT ^{1c}	INTERNAL AQUEOUS PHASE ^{2d}
80	50/40/10	40	Cetyl Dimethicone Copolyol ^{1e}	20% aqueous solution of a hair fixative resin (Polyquaternium-11) ^{2b}
81	70/20/10	20	Cetyl Dimethicone Copolyol ^{1e}	20% aqueous solution of a hair fixative resin (Polyquaternium-11) ^{2b}
82	70/20/10	20	Cetyl Dimethicone Copolyol ^{1e}	20% aqueous solution of a hair fixative resin (Polyquaternium-11) ^{2b}
83	70/20/10	20	Cetyl Dimethicone Copolyol ^{1e}	20% aqueous solution of a hair fixative resin (Polyquaternium-11) ^{2b}
84	80/17/3	17	Cyclomethicone and Dimethicone Copolyol Blend ^{4f}	20% aqueous solution of a hair fixative resin (Polyquaternium-11) ^{2b}
85	85/10/5	10	Dimethicone Copolyol ^{1g}	20% aqueous solution of a hair fixative resin (Polyquaternium-11) ^{2b}
86	85/10/5	10	Cetyl Dimethicone Copolyol ^{1e}	20% aqueous solution of a hair fixative resin (Polyquaternium-11) ^{2b}
87	87/10/3	10	Dimethicone Copolyol ^{1g}	20% aqueous solution of a hair fixative resin (Polyquaternium-11) ^{2b}
88	86.3/9.2/4.5	9.2	Cetyl Dimethicone Copolyol ^{1e}	20% aqueous solution of a hair fixative resin (Polyquaternium-11) ^{2b}
89	87/10/3	10	Dimethicone Copolyol ^{1g}	20% aqueous solution of a hair fixative resin (Polyquaternium-11) ^{2b}
90	78.9/14.6/6.5	14.6 ^{4g} (hydrocarbon)	Cetyl Dimethicone Copolyol ^{1e}	20% aqueous solution of a hair fixative resin (Polyquaternium-11) ^{2b}
91	80/12.6/7.4	12.6 ^{4g} (hydrocarbon)	Cetyl Dimethicone Copolyol ^{1e}	20% aqueous solution of a hair fixative resin (Polyquaternium-11) ^{2b}
92	80/12.6/7.4	12.6 ^{4g} (hydrocarbon)	Cetyl Dimethicone Copolyol ^{1e} (50%) and Laureth-4 (50%)	20% aqueous solution of a hair fixative resin (Polyquaternium-11) ^{2b}
93	80/10/10	10 ^m Cetyl Dimethicone ^m (nonvolatile silicone)	ABIL WE-09 ^m	D&C Red #40 Dye (2%)

TABLE III				
EXAMPLE	W/O/S ¹⁰	% SILICONE ²⁰	SURFACTANT ¹⁰	INTERNAL AQUEOUS PHASE ²⁰
94	82/11.6/6.4	11.6 ^m Cetyl Dimethicone ²⁰ (nonvolatile silicone)	Cetyl Dimethicone Copolyol ¹⁰	D&C Red #40 Dye (3%)
95	75/18/7	18 ^m Phenyl Trimethicone ²⁰ (nonvolatile silicone)	Cetyl Dimethicone Copolyol ¹⁰	Glycolic Acid (10%)

¹⁰ weight percent volatile silicone (cyclomethicone) or volatile hydrocarbon in the primary W/O emulsion;

²⁰ FLEXAN 130, hair fixative resin, available from National Starch & Chemical Corp., Bridgewater, NJ, available as a 30% active material;

²⁵ first topically-active compound incorporated into the internal aqueous phase of the primary emulsion

³⁰ GANTREZ[®] XL-80, hair fixative resin, available from ISP Chemicals, Inc., Wayne, NJ, added as a 5% active material;

³⁵ CROTEIN HKP, available from Croda, Inc., NY, NY, as a 50% active material;

⁴⁰ AJIDREW N-50, available from Ajinomoto U.S.A., Inc., Teaneck, NJ, as a 50% active material;

⁴⁵ DC Q2-5324, available from Dow Corning Corp., Midland, MI, as a 100% active material;

⁵⁰ AMERSIL ME358, available from Amerchol Corp., Edison, NJ, as a 100% active material;

⁵⁵ The cyclomethicone was replaced with a volatile hydrocarbon, PERMETHYL 101A, available from Presperse, Inc., South Plainfield, NJ;

⁶⁰ The cyclomethicone was replaced by a nonvolatile silicone;

⁶⁵ ABIL[®] Wax 9801, available as a 100% active material from Goldschmidt Chemical Co., Hopewell, VA;

⁷⁰ a blend of polyglyceryl-4 isostearate, cetyl dimethicone copolyol and hexyl laurate, available from Goldschmidt Chemical Co., Hopewell, VA; and

⁷⁵ DOW CORNING 556 FLUID, available as a 100% active material from Dow Corning Corp., Midland, MI.

Each W₁-O-W₂ multiple emulsion composition summarized in Table III was stable for a definite time period. The multiple emulsion compositions of Examples 42-95 were not optimized for long term stability. Therefore, the stability of the multiple emulsion compositions varied with the identity and amount of composition ingredients, especially the emulsifiers, and with the manufacturing conditions. For example, by using low shear and relatively low mixing speeds (e.g., about 100 to about 200 rpm on a bench-scale mixer), the external aqueous phase was able to emulsify the primary W₁/O emulsion. In every Example 42-95, the oil phase (O) of the primary emulsion isolated the first topically-active compound in the internal water phase (W₁) from the cationic hair conditioners present in the external aqueous phase (W₂).

The dye present in the compositions of Examples 61-68 and 79 dyed the hair after the cyclomethicone evaporated. The compositions of Examples 61-68 and 79 were non-staining to skin and equipment because the dyes are not present in the external aqueous phase. W₁/O primary emulsion droplets incorporating a water-soluble or water-dispersible dye was observed in magnified photographs of the compositions of Examples 61-65. Deposition of the dyes was observed

by a color change in the hair after application of the composition to the hair and hair drying. Deposition of the sunscreen benzophenone-4 of Example 66 was detected on a wool swatch by measurement of the absorbance of ultraviolet light.

The compositions of Examples 93-95 utilize a nonvolatile silicone compound as the oil phase. The first topically-active compound in each composition was released by rubbing the treated hair after the treated hair was rinsed.

As previously stated, two independent W₁-O-W₂ multiple emulsion compositions having different primary W₁/O emulsions can be combined without commingling of internal aqueous phases until after deposition and evaporation of the volatile oil phase. This was demonstrated in a test wherein two different primary emulsions were prepared. The first primary emulsion incorporated a red dye in the internal aqueous phase. The second primary emulsion incorporated a blue dye in the internal aqueous phase. Each primary emulsion was added to individual portions of an external aqueous phase to provide two independent W₁-O-W₂ multiple emulsion compositions. The two multiple emulsion compositions then were combined. The dye-containing droplets of the two primary emulsions were distinct and did not coalesce, commingle or transfer dyes when viewed through a microscope.

The following Examples 96-111 illustrate other primary W₁/O emulsions that have been prepared. Each of these primary emulsions can be incorporated into an external aqueous phase to provide a W₁-O-W₂ multiple emulsion composition of the present invention.

In Examples 96 through 111, the volatile oil phase was cyclomethicone (DOW CORNING 344 FLUID). The surfactant phase was a combination of laureth-1 and laureth-4. In preparing the compositions of Examples 96 through 111, the cyclomethicone, laureth-1 and laureth-4, in a weight ratio of 70:20:10, respectively, were admixed until homogeneous. The internal aqueous phase comprised water, the first topically-active compound and sugar. The aqueous phase was 80% by weight of the primary emulsion. The sugar was an optional ingredient included in the internal aqueous phase to match the refractive index of the aqueous phase to the refractive index of the volatile organic phase, and thereby provide a transparent primary W₁/O emulsion in most cases.

Admixing the aqueous phase with the solution of the volatile oil phase and the surfactant phase provided a clear, stable primary (W₁/O) emulsion. Table IV summarizes the final weight percent of each ingredient present in the primary emulsions of Examples 96 through 111, the final use of the composition, and the stability of the primary emulsion.

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TABLE IV

Ex.	Internal Aqueous Phase			Volatile Organic Phase			Surfactant Phase			Composition	
	% Sugar ¹¹⁾	First Topically-effective compound	% Topically-effective compound	% Water	% Cyclomethicone	% Laureth-1	% Laureth-4	Product Type	Stability		
96	36.4	Hyaluronic acid	0.13	44.1	13.6	3.88	1.94	Skin conditioner	stable		
97	29.0	Sodium salicylate	3.90	47.0	14.0	4.00	2.00	Skin conditioner	stable		
98	31.8	Urea	7.60	47.1	9.49	2.70	1.35	Skin conditioner	stable		
99	35.5	85% lactic acid	4.90	45.0	10.2	2.93	1.46	Skin conditioner	stable		
100	33.4	5% glycerin	3.60	46.8	11.3	3.24	1.62	Skin conditioner	stable		
101	32.2	Vitamin C	4.29	45.1	12.9	3.69	1.84	water-soluble vitamin	stable		
102	35.3	Vitamin B ₁	0.59	45.0	13.4	3.83	1.92	water-soluble vitamin	stable		
103	34.3	Pyridoxine hydrochloride	1.32	45.1	13.5	3.87	1.94	water-soluble vitamin	stable		
104	37.1	Panthenol	1.90	46.3	10.3	2.95	1.47	water-soluble vitamin	stable		
105	35.7	Nicotinic acid	0.42	44.5	13.6	3.88	1.94	cosmetic	stable		
106	35.7	Boric acid	1.08	43.8	13.7	3.90	1.95	medicament	stable		
107	34.9	Sodium sulfacetamide	2.74	48.5	9.70	2.77	1.39	bactericide	stable		
108	32.5	Triclosan	1.03	37.6	20.2	5.77	2.88	bactericide	stable		

Ex.	Internal Aqueous Phase			Volatile Organic Phase			Surfactant Phase			Composition	
	% Sugar ¹¹⁾	First Topically-effective compound	% Topically-effective compound	% Water	% Cyclomethicone	% Laureth-1	% Laureth-4	% Laureth-1	% Laureth-4	Product Type	Stability
109	37.6	Indometacin	1.90	43.4	12.8	2.91	1.43	anti-inflammatory agent	stable		
110	39.1	87% bisabolol	0.70	45.1	10.6	3.03	1.51	anti-irritant	stable		
111	31.9	Silver sulfadiazine	3.30	36.8	19.6	5.61	2.81	medicament	stable		

¹¹⁾ All % are by weight of the primary W/O emulsion.

Each composition of Examples 96-111 can be incorporated in a W₁-O-W₂ multiple emulsion composition of the present invention, wherein the external aqueous phase comprises a skin cleanser. The skin cleanser cleans the skin

and is removed from the skin during the rinsing step. The first topically-active compound remains on the skin to perform its intended function after evaporation of the cyclomethicone.

A W₁-O-W₂ multiple emulsion composition of the present invention demonstrates excellent esthetic and functional properties, such as the ability to treat the hair or skin with incompatible topically-active compounds from a single composition. The present multiple emulsion compositions also provide a more efficacious and economical delivery of water-soluble topically-active compounds from a rinse-off product. The compositions also are phase stable at room temperature for extended storage periods.

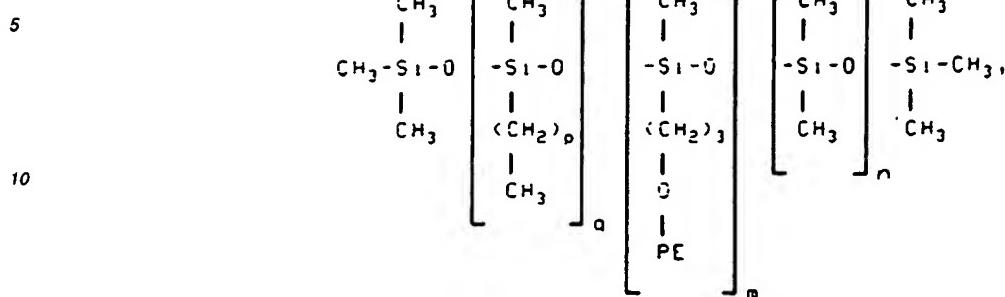
In addition, the present W₁-O-W₂ multiple emulsion compositions have the additional benefit of keeping a water-soluble, first topically-active compound in the internal aqueous phase from contacting individuals that apply the multiple emulsion composition, or from contacting workplace articles, such as sinks and countertops. The volatility of the oil phase of the primary emulsion also can be selected and adjusted for release of the first topically-active compound at a predetermined temperature range, e.g., at normal hair drying, at blow drying or at curling iron temperatures.

It should be understood that the foregoing detailed description is given merely by way of illustration. Obviously, many modifications and variations of the invention as hereinbefore set forth can be made without departing from the spirit and scope thereof and therefore only such limitations should be imposed as are indicated by the appended claims.

Claims

1. A rinse-off water-in-oil-in-water multiple emulsion composition comprising:
 - (a) about 40% to about 99% by weight of the composition of an external aqueous phase comprising water; and
 - (b) about 1% to about 60% by weight of the composition of a primary water-in-oil emulsion, said primary emulsion comprising:
 - (i) about 1% to about 95% by weight of the primary emulsion of an aqueous phase comprising a topically-effective amount of a first topically-active compound and water,
 - (ii) about 0.5% to about 95% by weight of the primary emulsion of an oil phase comprising a volatile silicone compound, a nonvolatile silicone compound, a volatile hydrocarbon compound, a nonvolatile hydrocarbon compound, or a mixture thereof, and
 - (iii) about 0.1% to about 20% by weight of the primary emulsion of a surfactant phase comprising a silicon-free surfactant or surfactant blend having an HLB value of about 10 or less, an oil-soluble silicon-based surfactant, an oil-soluble polymeric surfactant, or a mixture thereof.
2. The composition of claim 1 wherein the external aqueous phase further comprises 0% to about 15% by weight of the external phase of an emulsifier selected from the group consisting of an anionic surfactant, a cationic surfactant, an amphoteric surfactant, a water-soluble polymeric surfactant, a water-soluble silicon-containing surfactant, a non-ionic surfactant having an HLB value greater than about 10, a surfactant system capable of forming liquid crystals around droplets of the primary emulsion, and mixtures thereof; and 0% to about 30% by weight of the external phase of a second topically-active compound.
3. The composition of claim 1 wherein the external aqueous phase is free of a polymeric gelling agent.
4. The composition of claim 1 wherein the composition is a liquid having a viscosity of about 1 to about 15,000 centipoise.
- 45 5. The composition of claim 1 wherein the composition is a cream having a viscosity of about 50,000 to about 1,200,000 centipoise and the external aqueous phase is free of a polymeric gelling agent.
6. The composition of claim 1 wherein the primary emulsion is present in the form of droplets having a diameter of about 5 to about 1000 microns.
- 50 7. The composition of claim 2 wherein the first topically-active compound and the second topically-active compound are different compounds.
8. The composition of claim 2 wherein the first topically-active compound and the second topically-active compound are identical compounds.
- 55 9. The composition of claim 2 wherein the first topically-active compound and the second topically-active compound are incompatible compounds.

10. The composition of claim 1 wherein the aqueous phase of the primary emulsion is present in an amount of about 10% to about 95% by weight of the primary emulsion.
- 5 11. The composition of claim 1 wherein the first topically-active compound is present in an amount of about 0.1% to about 30% by weight of the primary emulsion.
12. The composition of claim 1 wherein the first topically-active compound is water soluble.
- 10 13. The composition of claim 1 wherein the first topically-active compound is selected from the group consisting of a hair conditioner, a skin conditioner, a hair cleanser, a skin cleanser, a hair fixative, a hair dye, a hair growth promoter, a deodorant, a skin care compound, a permanent wave compound, a hair relaxer, a hair straightener, an antibacterial compound, a topically-active drug, an antifungal compound, an anti-inflammatory compound, a topical anesthetic, a sunscreen, a dermatitis medication, an acne preparation, an oxidizing agent, an antioxidant, an optical brightener, a hair bleaching agent, a hair reducing agent, an ultraviolet light absorber, a sunburn reliever, a self-tanning compound, an anti-itch compound, and mixtures thereof.
- 15 14. The composition of claim 1 wherein the oil phase comprises a volatile silicone compound.
- 20 15. The composition of claim 14 wherein the cyclomethicone is selected from the group consisting of hexamethylcyclotrisiloxane, octamethylcyclotetrasiloxane, decamethylcyclopentasiloxane, dodecamethylcyclohexasiloxane, and mixtures thereof.
- 25 16. The composition of claim 1 wherein the oil phase comprises a volatile hydrocarbon compound having about 10 to about 30 carbon atoms.
- 30 17. The composition of claim 1 wherein the oil phase comprises a nonvolatile silicone compound.
18. The composition of claim 1 wherein the oil phase comprises a nonvolatile hydrocarbon compound.
- 35 19. The composition of claim 1 wherein the oil phase comprises a volatile oil phase and a nonvolatile oil phase.
20. The composition of claim 1 wherein the surfactant phase comprises a silicon-free surfactant or surfactant blend having an HLB value of about 1 to about 7.
- 35 21. The composition of claim 1 wherein the surfactant phase comprises one silicon-free surfactant having an HLB value of about 0.1 to about 10.
- 40 22. The composition of claim 1 wherein the surfactant phase comprises a silicon-free surfactant blend having an HLB value of about 1 to about 10, said surfactant blend comprising a first surfactant having an HLB value of about 0.1 to about 10 and a second surfactant having an HLB greater than about 10.
- 45 23. The composition of claim 1 wherein the surfactant phase comprises a silicon-free nonionic surfactant selected from the group consisting of a polyoxyethylene ether of a fatty (C_6-C_{22}) alcohol, a polyoxyethylene/polyoxypropylene ether of a fatty (C_6-C_{22}) alcohol, an ethoxylated alkylphenol, a polyethylene glycol ether of methyl glucose, a polyethylene ether of sorbitol, and mixtures thereof.
24. The composition of claim 1 wherein the surfactant phase comprises an oil-soluble silicon-based surfactant.
- 50 25. The composition of claim 24 wherein the silicon-based surfactant comprises a dimethicone copolyol having about 15 or less total monomer units of ethylene oxide, propylene oxide or a mixture thereof, an alkyl dimethicone copolyol having the structure:



wherein p is a numeral from 7 through 17;

q is a numeral from 1 through 100;

m is a numeral from 1 through 40;

n is a numeral from 0 through 200; and

20 PE is $(\text{C}_2\text{H}_4\text{O})_a(\text{C}_3\text{H}_6\text{O})_b\text{-H}$ having a molecular weight of about 250 to about 2000, wherein a and b are selected such that the weight ratio of $\text{C}_2\text{H}_4\text{O}/\text{C}_3\text{H}_6\text{O}$ is from 100/0 to 20/80, or a mixture thereof.

- 25 26. The composition of claim 1 wherein the surfactant phase comprises an oil-soluble polymeric surfactant capable of forming water-in-oil emulsions and having repeating ethoxy units, propoxy units, butoxy units or combinations thereof, and includes less than 40% by weight ethoxy units.
- 30 27. The composition of claim 1 wherein the surfactant phase comprises an oil-soluble silicon-based surfactant, and the oil phase comprises a volatile silicone compound or a nonvolatile silicone compound.
- 35 28. The composition of claim 1 wherein the surfactant phase comprises a silicon-free nonionic surfactant or an oil-soluble polymeric surfactant and the oil phase comprises a volatile hydrocarbon compound or nonvolatile hydrocarbon compound.
- 40 29. The composition of claim 1 wherein the oil phase comprises a combination of a silicone compound and hydrocarbon compound, and the surfactant phase comprises a combination of a silicon-based surfactant and a silicon-free non-ionic surfactant, an oil-soluble polymeric surfactant or a mixture thereof.
- 30 30. The composition of claim 2 wherein the external aqueous phase comprises about 0.1% to about 10% by weight of the external aqueous phase of an emulsifier.
- 45 31. The composition of claim 2 wherein the nonionic surfactant has an HLB value of at least about 12.
32. The composition of claim 2 wherein the water-soluble polymeric surfactant has repeating ethoxy units, propoxy units, butoxy units or combinations thereof, and includes 20% by weight or more ethoxy units.
- 50 33. The composition of claim 2 wherein the water-soluble silicon-containing surfactant is a dimethicone copolyol having about 15 or more monomer units of ethylene oxide, propylene oxide or a mixture thereof.
34. The composition of claim 2 wherein the surfactant system capable of forming liquid crystals is selected from the group consisting of a quaternary ammonium compound having two long alkyl ($\text{C}_8\text{-C}_{22}$) chains, a di-long chain alkyl ($\text{C}_8\text{-C}_{22}$) amine, a long chain fatty ($\text{C}_8\text{-C}_{22}$) alcohol, an ethoxylated fatty ($\text{C}_8\text{-C}_{22}$) alcohol, a double-tailed anionic surfactant, and a double-tailed phospholipid, and mixtures thereof.
- 55 35. The composition of claim 2 wherein the first topically-active compound comprises a hair fixative resin, a hair reducing agent, a protein, an oxidizing agent, a water-soluble hair dye, a sunscreen, a humectant, or an optical brightener, and the second topically-active compound comprises a cationic hair conditioner or an anionic hair cleanser.

36. The composition of claim 2 wherein the first topically-active compound comprises a cationic hair conditioner, a deodorant, a skin care product, a bacteriocide, a topical medicament or a mixture thereof, and the second topically-active compound comprises an anionic cleanser.
- 5 37. The composition of claim 2 wherein the first topically-active compound comprises a neutralizer for reducing agents and the second topically-active compound comprises a reducing agent for permanent hair waving, cysteine, cysteamine, or a hair conditioner.
- 10 38. The composition of claim 2 wherein the first topically-active compound comprises a reducing agent for permanent hair waving, and the second topically-active compound comprises a hair cleanser or a hair conditioner.
- 15 39. The composition of claim 2 wherein the first topically-active compound comprises a hair conditioner, and the second topically-active compound comprises an oxidizing agent.
- 20 40. The composition of claim 2 wherein the first topically-active compound comprises a skin conditioner and the second topically-active compound comprises a make-up remover.
- 25 41. The composition of claim 2 wherein the first topically-active compound comprises a sunburn reliever, a sunscreen, a skin conditioner, a bacteriocide, a deodorant, an antifungal compound, anesthetic, an anti-inflammatory, an anti-irritant, an anti-itch compound, a medicament, a self-tanning compound, a dermatitis medication, or a mixture thereof, and the second topically-active compound comprises a skin cleanser.
- 30 42. The composition of claim 2 wherein the first topically-active compound comprises a self-tanning compound, and the second topically-active compound comprises a skin conditioner.
- 35 43. The composition of claim 2 wherein the first topically-active compound comprises a hair conditioner, a hair dye, a hair growth promoter, a hair bleaching agent, or a mixture thereof, and the second topically-active compound comprises a hair cleanser.
- 40 44. The composition of claim 2 wherein the first topically-active compound comprises a hair dye, and the second topically-active compound comprises a hair conditioner, a bleaching agent, or a mixture thereof.
- 45 45. The composition of claim 2 wherein the first topically-active compound comprises a water-soluble hair conditioner and the second topically-active compound comprises a substantive hair conditioner.
- 50 46. The composition of claim 2 wherein the first topically-active compound comprises a sunscreen, a protein, a humectant, or a mixture thereof, and the second topically-active compound comprises a hair conditioner.
- 55 47. The composition of claim 2 wherein the hair fixative resin is selected from the group consisting of a polyvinylpyrrolidone, Polyquaternium-11, PVP/dimethylaminoethylmethacrylate copolymer, a sodium polystyrene sulfonate, a polyethylene glycol, and mixtures thereof.
- 60 48. The composition of claim 1 wherein the surfactant phase comprises cetyl dimethicone copolyol, dimethicone copolyol, stearoxy dimethicone, laureth-4, or a mixture thereof, and the oil phase comprises a volatile silicone compound.
- 65 49. A rinse-off, water-in-oil-in-water multiple emulsion composition comprising:
- 70 (a) about 50% to about 95% by weight of the composition of an external aqueous phase, said external aqueous phase comprising:
- 75 (i) about 0.1% to about 30% by weight of the external aqueous phase of a second topically-active compound;
- 80 (ii) about 0.1% to about 10% by weight of the external aqueous phase of an emulsifier, selected from the group consisting of an anionic surfactant, a cationic surfactant, an amphoteric surfactant, a water-soluble polymeric surfactant, a surfactant system capable of forming liquid crystals, a nonionic surfactant or surfactant blend having an HLB value of greater than about 10, and mixtures thereof; and
- 85 (iii) water; and
- 90 (b) about 5% to about 50% by weight of the composition of a primary water-in-oil emulsion, said primary emulsion comprising:

(i) about 10% to about 95% by weight of the primary emulsion of an aqueous phase, said aqueous phase comprising (A) water and (B) about 0.1% to about 30% by weight of the primary emulsion of a first topically-active compound;

5 (ii) about 0.5% to about 80% by weight of the primary emulsion of an oil phase comprising a volatile silicone compound, a volatile hydrocarbon compound, or a mixture thereof; and

(iii) about 0.1% to about 15% by weight of the primary emulsion of a surfactant phase comprising a silicon-free surfactant or surfactant blend having an HLB value of about 1 to about 7, an oil-soluble silicon-based surfactant, an oil-soluble polymeric surfactant, or a mixture thereof.

10 50. A rinse-off, water-in-oil-in-water multiple emulsion composition comprising:

(a) about 40% to about 99% by weight of the composition of an external aqueous phase comprising water; and
 (b) about 1% to about 60% by weight of the composition of a primary emulsion, said primary emulsion comprising:

15 (i) about 0.01% to about 57% by weight of the composition of an aqueous phase comprising water and about 0.001% to about 17.1% by weight of the composition of a first topically-active compound,

(ii) about 0.005% to about 57% by weight of the composition of an oil phase comprising a volatile silicone compound, a nonvolatile silicone compound, a volatile hydrocarbon compound, a nonvolatile hydrocarbon compound, or a mixture thereof, and

20 (iii) about 0.001% to about 12% by weight of the composition of a surfactant phase comprising a silicon-free surfactant or surfactant blend having an HLB value of about 10 or less, an oil-soluble silicon-based surfactant, an oil-soluble polymeric surfactant, or a mixture thereof.

25 51. The composition of claim 50 wherein the external phase further comprises 0% to about 14.85% by weight of the composition of an emulsifier, and 0% to about 30% by weight of the composition of a second topically-active compound.

30 52. A method of treating the skin or hair with a topically-active compound comprising topically applying an effective amount of a water-in-oil-in-water multiple emulsion composition to skin or hair, said composition comprising:

(a) about 40% to about 99% by weight of the composition of an external aqueous phase comprising water; and
 (b) about 1% to about 60% by weight of the composition of a primary water-in-oil emulsion, said primary emulsion comprising:

35 (i) about 1% to about 95% by weight of the primary emulsion of an aqueous phase comprising a topically-effective amount of a first topically-active compound and water,

(ii) about 0.5% to about 95% by weight of the primary emulsion of an oil phase comprising a volatile silicone compound, a nonvolatile silicone compound, a volatile hydrocarbon compound, a nonvolatile hydrocarbon compound, or a mixture thereof, and

40 (iii) about 0.1% to about 20% by weight of the primary emulsion of a surfactant phase comprising a silicon-free surfactant or surfactant blend having an HLB value of about 10 or less, an oil-soluble silicon-based surfactant, an oil-soluble polymeric surfactant, or a mixture thereof;

45 then rinsing the composition from the skin or hair, wherein droplets of the primary emulsion remain in contact with the skin or hair to release the first topically-active compound.